

(predictive) purposes to thereby treat an individual prophylactically. Accordingly, one aspect of the present invention relates to diagnostic assays for determining INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 protein and/or nucleic acid expression as well as INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 activity, in the context of a biological sample (*e.g.*, blood, serum, cells, tissue) to thereby determine whether an individual is afflicted with a disease or disorder, or is at risk of developing a disorder, associated with aberrant or unwanted INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 gene expression or activity. The invention also provides for prognostic (or predictive) assays for determining whether an individual is at risk of developing a disorder associated with INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 protein or nucleic acid expression or activity. For example, mutations in a gene can be assayed in a biological sample. Such assays can be used for prognostic or predictive purpose to thereby prophylactically treat an individual prior to the onset of a disorder characterized by or associated with protein or nucleic acid expression or activity.

As an alternative to making determinations based on the absolute expression level of selected genes, determinations may be based on the normalized expression levels of these genes. Expression levels are normalized by correcting the absolute expression level of a INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 gene by comparing its expression to the expression of a gene that is not a INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378, *e.g.*, a housekeeping gene that is constitutively expressed. Suitable genes for normalization include housekeeping genes such as the actin gene. This normalization allows the comparison of the expression level in one sample, *e.g.*, a patient sample, to another sample, *e.g.*, a non-disease sample, or between samples from different sources.

Alternatively, the expression level can be provided as a relative expression level. To determine a relative expression level of a gene, the level of expression of the gene is determined for 10 or more samples of different cell isolates, preferably 50 or more samples, prior to the determination of the expression level for the sample in question. The mean expression level of each of the genes assayed in the larger number of samples is determined and this is used as a baseline expression level for the gene(s) in question. The expression level of the gene determined for the test sample (absolute level of expression) is then divided by the mean expression value obtained for that gene. This provides a relative expression level and aids in identifying extreme cases of disease.

Preferably, the samples used in the baseline determination will be from diseased or from non-diseased cells of tissue. The choice of the cell source is dependent on the use of

the relative expression level. Using expression found in normal tissues as a mean expression score aids in validating whether the INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 gene assayed is diseased cell-type specific (versus normal cells). Such a use is particularly important in identifying whether a INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295,
5 TANGO 354, or TANGO 378 gene can serve as a target gene. In addition, as more data is accumulated, the mean expression value can be revised, providing improved relative expression values based on accumulated data. Expression data from cells provide a means for grading the severity of the disease state.

Another aspect of the invention pertains to monitoring the influence of agents (*e.g.*,
10 drugs, compounds) on the expression or activity of INTERCEPT 340, MANGO 003, MANGO 347, TANGO 272, TANGO 295, TANGO 354, or TANGO 378 genes in clinical trials.

These and other agents are described in further detail in the following sections.

15 1. Diagnostic Assays

An exemplary method for detecting the presence or absence of a polypeptide or nucleic acid of the invention in a biological sample involves obtaining a biological sample from a test subject and contacting the biological sample with a compound or an agent capable of detecting a polypeptide or nucleic acid (*e.g.*, mRNA, genomic DNA) of the
20 invention such that the presence of a polypeptide or nucleic acid of the invention is detected in the biological sample. A preferred agent for detecting mRNA or genomic DNA encoding a polypeptide of the invention is a labeled nucleic acid probe capable of hybridizing to mRNA or genomic DNA encoding a polypeptide of the invention. The nucleic acid probe can be, for example, a full-length cDNA, such as the nucleic acid of SEQ
25 ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, 19, 21, 22, 24, 25, 27, 28 or 30, or a portion thereof, such as an oligonucleotide of at least 15, 30, 50, 100, 250 or 500 nucleotides in length and sufficient to specifically hybridize under stringent conditions to a mRNA or genomic DNA encoding a polypeptide of the invention. Other suitable probes for use in the diagnostic assays of the invention are described herein.

30 A preferred agent for detecting a polypeptide of the invention is an antibody capable of binding to a polypeptide of the invention, preferably an antibody with a detectable label. Antibodies can be polyclonal, or more preferably, monoclonal. An intact antibody, or a fragment thereof (*e.g.*, Fab or F(ab')₂) can be used. The term "labeled", with regard to the probe or antibody, is intended to encompass direct labeling of the probe or antibody by
35 coupling (*i.e.*, physically linking) a detectable substance to the probe or antibody, as well as indirect labeling of the probe or antibody by reactivity with another reagent that is directly

labeled. Examples of indirect labeling include detection of a primary antibody using a fluorescently labeled secondary antibody and end-labeling of a DNA probe with biotin such that it can be detected with fluorescently labeled streptavidin. The term "biological sample" is intended to include tissues, cells and biological fluids isolated from a subject, as well as tissues, cells and fluids present within a subject. That is, the detection method of the invention can be used to detect mRNA, protein, or genomic DNA in a biological sample *in vitro* as well as *in vivo*. For example, *in vitro* techniques for detection of mRNA include Northern hybridizations and *in situ* hybridizations. *In vitro* techniques for detection of a polypeptide of the invention include enzyme linked immunosorbent assays (ELISAs), Western blots, immunoprecipitations and immunofluorescence. *In vitro* techniques for detection of genomic DNA include Southern hybridizations. Furthermore, *in vivo* techniques for detection of a polypeptide of the invention include introducing into a subject a labeled antibody directed against the polypeptide. For example, the antibody can be labeled with a radioactive marker whose presence and location in a subject can be detected by standard imaging techniques.

In one embodiment, the biological sample contains protein molecules from the test subject. Alternatively, the biological sample can contain mRNA molecules from the test subject or genomic DNA molecules from the test subject. A preferred biological sample is a peripheral blood leukocyte sample isolated by conventional means from a subject.

In another embodiment, the methods further involve obtaining a control biological sample from a control subject, contacting the control sample with a compound or agent capable of detecting a polypeptide of the invention or mRNA or genomic DNA encoding a polypeptide of the invention, such that the presence of the polypeptide or mRNA or genomic DNA encoding the polypeptide is detected in the biological sample, and comparing the presence of the polypeptide or mRNA or genomic DNA encoding the polypeptide in the control sample with the presence of the polypeptide or mRNA or genomic DNA encoding the polypeptide in the test sample.

The invention also encompasses kits for detecting the presence of a polypeptide or nucleic acid of the invention in a biological sample (a test sample). Such kits can be used to determine if a subject is suffering from or is at increased risk of developing a disorder associated with aberrant expression of a polypeptide of the invention (*e.g.*, a proliferative disorder, *e.g.*, psoriasis or cancer). For example, the kit can comprise a labeled compound or agent capable of detecting the polypeptide or mRNA encoding the polypeptide in a biological sample and means for determining the amount of the polypeptide or mRNA in the sample (*e.g.*, an antibody which binds the polypeptide or an oligonucleotide probe which binds to DNA or mRNA encoding the polypeptide). Kits can also include instructions for observing that the tested subject is suffering from or is at risk of developing

a disorder associated with aberrant expression of the polypeptide if the amount of the polypeptide or mRNA encoding the polypeptide is above or below a normal level.

For antibody-based kits, the kit can comprise, for example: (1) a first antibody (*e.g.*, attached to a solid support) which binds to a polypeptide of the invention; and, optionally, (2) a second, different antibody which binds to either the polypeptide or the first antibody and is conjugated to a detectable agent.

For oligonucleotide-based kits, the kit can comprise, for example: (1) an oligonucleotide, *e.g.*, a detectably labeled oligonucleotide, which hybridizes to a nucleic acid sequence encoding a polypeptide of the invention or (2) a pair of primers useful for amplifying a nucleic acid molecule encoding a polypeptide of the invention. The kit can also comprise, *e.g.*, a buffering agent, a preservative, or a protein stabilizing agent. The kit can also comprise components necessary for detecting the detectable agent (*e.g.*, an enzyme or a substrate). The kit can also contain a control sample or a series of control samples which can be assayed and compared to the test sample contained. Each component of the kit is usually enclosed within an individual container and all of the various containers are within a single package along with instructions for observing whether the tested subject is suffering from or is at risk of developing a disorder associated with aberrant expression of the polypeptide.

2. Prognostic Assays

The methods described herein can furthermore be utilized as diagnostic or prognostic assays to identify subjects having or at risk of developing a disease or disorder associated with aberrant expression or activity of a polypeptide of the invention. For example, the assays described herein, such as the preceding diagnostic assays or the following assays, can be utilized to identify a subject having or at risk of developing a disorder associated with aberrant expression or activity of a polypeptide of the invention. Alternatively, the prognostic assays can be utilized to identify a subject having or at risk for developing such a disease or disorder. Thus, the present invention provides a method in which a test sample is obtained from a subject and a polypeptide or nucleic acid (*e.g.*, mRNA, genomic DNA) of the invention is detected, wherein the presence of the polypeptide or nucleic acid is diagnostic for a subject having or at risk of developing a disease or disorder associated with aberrant expression or activity of the polypeptide. As used herein, a "test sample" refers to a biological sample obtained from a subject of interest. For example, a test sample can be a biological fluid (*e.g.*, serum), cell sample, or tissue.

Furthermore, the prognostic assays described herein can be used to determine whether a subject can be administered an agent (*e.g.*, an agonist, antagonist, peptidomimetic, protein, peptide, nucleic acid, small molecule, or other drug candidate) to

treat a disease or disorder associated with aberrant expression or activity of a polypeptide of the invention. For example, such methods can be used to determine whether a subject can be effectively treated with a specific agent or class of agents (e.g., agents of a type which decrease activity of the polypeptide). Thus, the present invention provides methods for determining whether a subject can be effectively treated with an agent for a disorder
5 associated with aberrant expression or activity of a polypeptide of the invention in which a test sample is obtained and the polypeptide or nucleic acid encoding the polypeptide is detected (e.g., wherein the presence of the polypeptide or nucleic acid is diagnostic for a subject that can be administered the agent to treat a disorder associated with aberrant expression or activity of the polypeptide).

10 The methods of the invention can also be used to detect genetic lesions or mutations in a gene of the invention, thereby determining if a subject with the lesioned gene is at risk for a disorder characterized aberrant expression or activity of a polypeptide of the invention. In preferred embodiments, the methods include detecting, in a sample of cells from the subject, the presence or absence of a genetic lesion or mutation characterized by at least one
15 of an alteration affecting the integrity of a gene encoding the polypeptide of the invention, or the mis-expression of the gene encoding the polypeptide of the invention. For example, such genetic lesions or mutations can be detected by ascertaining the existence of at least one of: 1) a deletion of one or more nucleotides from the gene; 2) an addition of one or more nucleotides to the gene; 3) a substitution of one or more nucleotides of the gene; 4) a
20 chromosomal rearrangement of the gene; 5) an alteration in the level of a messenger RNA transcript of the gene; 6) an aberrant modification of the gene, such as of the methylation pattern of the genomic DNA; 7) the presence of a non-wild type splicing pattern of a messenger RNA transcript of the gene; 8) a non-wild type level of a the protein encoded by the gene; 9) an allelic loss of the gene; and 10) an inappropriate post-translational
25 modification of the protein encoded by the gene. As described herein, there are a large number of assay techniques known in the art which can be used for detecting lesions in a gene.

In certain embodiments, detection of the lesion involves the use of a probe/primer in a polymerase chain reaction (PCR) (see, e.g., U.S. Patent NOs. 4,683,195 and 4,683,202),
30 such as anchor PCR or RACE PCR, or, alternatively, in a ligation chain reaction (LCR) (see, e.g., Landegran et al., 1988, *Science* 241:1077-80; and Nakazawa et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:360-4), the latter of which can be particularly useful for detecting point mutations in a gene (see, e.g., Abravaya et al., 1995, *Nucleic Acids Res.* 23:675-82). This method can include the steps of collecting a sample of cells from a patient, isolating
35 nucleic acid (e.g., genomic, mRNA or both) from the cells of the sample, contacting the nucleic acid sample with one or more primers which specifically hybridize to the selected

gene under conditions such that hybridization and amplification of the gene (if present) occurs, and detecting the presence or absence of an amplification product, or detecting the size of the amplification product and comparing the length to a control sample. It is anticipated that PCR and/or LCR may be desirable to use as a preliminary amplification step in conjunction with any of the techniques used for detecting mutations described
5 herein.

Alternative amplification methods include: self sustained sequence replication (Guatelli et al., 1990, *Proc. Natl. Acad. Sci. USA* 87:1874-78), transcriptional amplification system (Kwoh, et al., 1989, *Proc. Natl. Acad. Sci. USA* 86:1173-7), Q-Beta Replicase (Lizardi et al., 1988, *Bio/Technology* 6:1197), or any other nucleic acid amplification
10 method, followed by the detection of the amplified molecules using techniques well known to those of skill in the art. These detection schemes are especially useful for the detection of nucleic acid molecules if such molecules are present in very low numbers.

In an alternative embodiment, mutations in a selected gene from a sample cell can be identified by alterations in restriction enzyme cleavage patterns. For example, sample
15 and control DNA is isolated, amplified (optionally), digested with one or more restriction endonucleases, and fragment length sizes are determined by gel electrophoresis and compared. Differences in fragment length sizes between sample and control DNA indicates mutations in the sample DNA. Moreover, the use of sequence specific ribozymes (*see, e.g.*, U.S. Patent No. 5,498,531) can be used to score for the presence of specific mutations by
20 development or loss of a ribozyme cleavage site.

In other embodiments, genetic mutations can be identified by hybridizing a sample and control nucleic acids, *e.g.*, DNA or RNA, to high density arrays containing hundreds or thousands of oligonucleotides probes (Cronin et al., 1996, *Human Mutation* 7:244-55; Kozal et al., 1996, *Nature Medicine* 2:753-9). For example, genetic mutations can be
25 identified in two-dimensional arrays containing light-generated DNA probes as described in Cronin et al., *supra*. Briefly, a first hybridization array of probes can be used to scan through long stretches of DNA in a sample and control to identify base changes between the sequences by making linear arrays of sequential overlapping probes. This step allows the identification of point mutations. This step is followed by a second hybridization array that
30 allows the characterization of specific mutations by using smaller, specialized probe arrays complementary to all variants or mutations detected. Each mutation array is composed of parallel probe sets, one complementary to the wild-type gene and the other complementary to the mutant gene.

In yet another embodiment, any of a variety of sequencing reactions known in the
35 art can be used to directly sequence the selected gene and detect mutations by comparing the sequence of the sample nucleic acids with the corresponding wild-type (control)

sequence. Examples of sequencing reactions include those based on techniques developed by Maxim and Gilbert (1977, *Proc. Natl. Acad. Sci. USA* 74:560) or Sanger (1977, *Proc. Natl. Acad. Sci. USA* 74:5463). It is also contemplated that any of a variety of automated sequencing procedures can be utilized when performing the diagnostic assays developed by Naeve et al. (1995, *Bio/Techniques* 19:448-53), including sequencing by mass spectrometry (see, e.g., PCT Publication No. WO 94/16101; Cohen et al., 1996, *Adv. Chromatogr.* 36:127-62; and Griffin et al., 1993, *Appl. Biochem. Biotechnol.* 38:147-59).

Other methods for detecting mutations in a selected gene include methods in which protection from cleavage agents is used to detect mismatched bases in RNA/RNA or RNA/DNA heteroduplexes (Myers et al., 1985, *Science* 230:1242). In general, the technique of mismatch cleavage entails providing heteroduplexes formed by hybridizing (labeled) RNA or DNA containing the wild-type sequence with potentially mutant RNA or DNA obtained from a tissue sample. The double-stranded duplexes are treated with an agent which cleaves single-stranded regions of the duplex such as which will exist due to basepair mismatches between the control and sample strands. RNA/DNA duplexes can be treated with RNase to digest mismatched regions, and DNA/DNA hybrids can be treated with S1 nuclease to digest mismatched regions.

In other embodiments, either DNA/DNA or RNA/DNA duplexes can be treated with hydroxylamine or osmium tetroxide and with piperidine in order to digest mismatched regions. After digestion of the mismatched regions, the resulting material is then separated by size on denaturing polyacrylamide gels to determine the site of mutation. See, e.g., Cotton et al., 1988, *Proc. Natl. Acad. Sci. USA* 85:4397; Saleeba et al., 1992, *Methods Enzymol.* 217:286-95. In a preferred embodiment, the control DNA or RNA can be labeled for detection.

In still another embodiment, the mismatch cleavage reaction employs one or more proteins that recognize mismatched base pairs in double-stranded DNA (so called DNA mismatch repair enzymes) in defined systems for detecting and mapping point mutations in cDNAs obtained from samples of cells. For example, the mutY enzyme of *E. coli* cleaves A at G/A mismatches and the thymidine DNA glycosylase from HeLa cells cleaves T at G/T mismatches (Hsu et al., 1994, *Carcinogenesis* 15:1657-62). According to an exemplary embodiment, a probe based on a selected sequence, e.g., a wild-type sequence, is hybridized to a cDNA or other DNA product from a test cell(s). The duplex is treated with a DNA mismatch repair enzyme, and the cleavage products, if any, can be detected from electrophoresis protocols or the like. See, e.g., U.S. Patent No. 5,459,039.

In other embodiments, alterations in electrophoretic mobility will be used to identify mutations in genes. For example, single strand conformation polymorphism (SSCP) may be used to detect differences in electrophoretic mobility between mutant and wild type

nucleic acids (Orita et al., 1989, *Proc. Natl. Acad. Sci. USA* 86:2766; see also Cotton, 1993, *Mutat. Res.* 285:125-44; Hayashi, 1992, *Genet. Anal. Tech. Appl.* 9:73-9). Single-stranded DNA fragments of sample and control nucleic acids will be denatured and allowed to renature. The secondary structure of single-stranded nucleic acids varies according to sequence, and the resulting alteration in electrophoretic mobility enables the detection of even a single base change. The DNA fragments may be labeled or detected with labeled probes. The sensitivity of the assay may be enhanced by using RNA (rather than DNA), in which the secondary structure is more sensitive to a change in sequence. In a preferred embodiment, the subject method utilizes heteroduplex analysis to separate double stranded heteroduplex molecules on the basis of changes in electrophoretic mobility (Keen et al., 1991, *Trends Genet.* 7:5).

In yet another embodiment, the movement of mutant or wild-type fragments in polyacrylamide gels containing a gradient of denaturant is assayed using denaturing gradient gel electrophoresis (DGGE) (Myers et al., 1985, *Nature* 313:495). When DGGE is used as the method of analysis, DNA will be modified to insure that it does not completely denature, for example by adding a 'GC clamp of approximately 40 bp of high-melting GC-rich DNA by PCR. In a further embodiment, a temperature gradient is used in place of a denaturing gradient to identify differences in the mobility of control and sample DNA (Rosenbaum and Reissner, 1987, *Biophys. Chem.* 265:12753).

Examples of other techniques for detecting point mutations include, but are not limited to, selective oligonucleotide hybridization, selective amplification, or selective primer extension. For example, oligonucleotide primers may be prepared in which the known mutation is placed centrally and then hybridized to target DNA under conditions which permit hybridization only if a perfect match is found (Saiki et al., 1986, *Nature* 324:163; Saiki et al., 1989, *Proc. Natl. Acad. Sci. USA* 86:6230). Such allele specific oligonucleotides are hybridized to PCR amplified target DNA or a number of different mutations when the oligonucleotides are attached to the hybridizing membrane and hybridized with labeled target DNA.

Alternatively, allele specific amplification technology which depends on selective PCR amplification may be used in conjunction with the instant invention. Oligonucleotides used as primers for specific amplification may carry the mutation of interest in the center of the molecule (so that amplification depends on differential hybridization; Gibbs et al., 1989, *Nucleic Acids Res.* 17:2437-48) or at the extreme 3' end of one primer where, under appropriate conditions, mismatch can prevent or reduce polymerase extension (Prossner, 1993, *Tibtech* 11:238). In addition, it may be desirable to introduce a novel restriction site in the region of the mutation to create cleavage-based detection (Gasparini et al., 1992, *Mol. Cell Probes* 6:1). It is anticipated that in certain embodiments amplification may also be

performed using Taq ligase for amplification (Barany, 1991, *Proc. Natl. Acad. Sci. USA* 88:189). In such cases, ligation will occur only if there is a perfect match at the 3' end of the 5' sequence making it possible to detect the presence of a known mutation at a specific site by looking for the presence or absence of amplification.

5 The methods described herein may be performed, for example, by utilizing pre-packaged diagnostic kits comprising at least one probe nucleic acid or antibody reagent described herein, which may be conveniently used, *e.g.*, in clinical settings to diagnose patients exhibiting symptoms or family history of a disease or illness involving a gene encoding a polypeptide of the invention. Furthermore, any cell type or tissue, preferably peripheral blood leukocytes, in which the polypeptide of the invention is expressed may be
10 utilized in the prognostic assays described herein.

3. Pharmacogenomics

Agents, or modulators which have a stimulatory or inhibitory effect on activity or expression of a polypeptide of the invention as identified by a screening assay described
15 herein can be administered to individuals to treat (prophylactically or therapeutically) disorders associated with aberrant activity of the polypeptide. In conjunction with such treatment, the pharmacogenomics (*i.e.*, the study of the relationship between an individual's genotype and that individual's response to a foreign compound or drug) of the individual may be considered. Differences in metabolism of therapeutics can lead to severe toxicity or
20 therapeutic failure by altering the relation between dose and blood concentration of the pharmacologically active drug. Thus, the pharmacogenomics of the individual permits the selection of effective agents (*e.g.*, drugs) for prophylactic or therapeutic treatments based on a consideration of the individual's genotype. Such pharmacogenomics can further be used to determine appropriate dosages and therapeutic regimens. Accordingly, the activity of a
25 polypeptide of the invention, expression of a nucleic acid of the invention, or mutation content of a gene of the invention in an individual can be determined to thereby select appropriate agent(s) for therapeutic or prophylactic treatment of the individual.

Pharmacogenomics deals with clinically significant hereditary variations in the response to drugs due to altered drug disposition and abnormal action in affected persons.
30 *See, e.g.*, Linder, 1997, *Clin. Chem.* 43(2):254-66. In general, two types of pharmacogenetic conditions can be differentiated. Genetic conditions transmitted as a single factor altering the way drugs act on the body are referred to as "altered drug action." Genetic conditions transmitted as single factors altering the way the body acts on drugs are referred to as "altered drug metabolism". These pharmacogenetic conditions can occur
35 either as rare defects or as polymorphisms. For example, glucose-6-phosphate dehydrogenase deficiency (G6PD) is a common inherited enzymopathy in which the main

clinical complication is haemolysis after ingestion of oxidant drugs (anti-malarials, sulfonamides, analgesics, nitrofurans) and consumption of fava beans.

As an illustrative embodiment, the activity of drug metabolizing enzymes is a major determinant of both the intensity and duration of drug action. The discovery of genetic polymorphisms of drug metabolizing enzymes (e.g., N-acetyltransferase 2 (NAT 2) and cytochrome P450 enzymes CYP2D6 and CYP2C19) has provided an explanation as to why some patients do not obtain the expected drug effects or show exaggerated drug response and serious toxicity after taking the standard and safe dose of a drug. These polymorphisms are expressed in two phenotypes in the population, the extensive metabolizer (EM) and poor metabolizer (PM). The prevalence of PM is different among different populations. For example, the gene coding for CYP2D6 is highly polymorphic and several mutations have been identified in PM, which all lead to the absence of functional CYP2D6. Poor metabolizers of CYP2D6 and CYP2C19 quite frequently experience exaggerated drug response and side effects when they receive standard doses. If a metabolite is the active therapeutic moiety, a PM will show no therapeutic response, as demonstrated for the analgesic effect of codeine mediated by its CYP2D6-formed metabolite morphine. The other extreme are the so called ultra-rapid metabolizers who do not respond to standard doses. Recently, the molecular basis of ultra-rapid metabolism has been identified to be due to CYP2D6 gene amplification.

Thus, the activity of a polypeptide of the invention, expression of a nucleic acid encoding the polypeptide, or mutation content of a gene encoding the polypeptide in an individual can be determined to thereby select appropriate agent(s) for therapeutic or prophylactic treatment of the individual. In addition, pharmacogenetic studies can be used to apply genotyping of polymorphic alleles encoding drug-metabolizing enzymes to the identification of an individual's drug responsiveness phenotype. This knowledge, when applied to dosing or drug selection, can avoid adverse reactions or therapeutic failure and thus enhance therapeutic or prophylactic efficiency when treating a subject with a modulator of activity or expression of the polypeptide, such as a modulator identified by one of the exemplary screening assays described herein.

4. Monitoring of Effects During Clinical Trials

Monitoring the influence of agents (e.g., drugs, compounds) on the expression or activity of a polypeptide of the invention (e.g., the ability to modulate aberrant cell proliferation chemotaxis, and/or differentiation) can be applied not only in basic drug screening, but also in clinical trials. For example, the effectiveness of an agent, as determined by a screening assay as described herein, to increase gene expression, protein levels or protein activity, can be monitored in clinical trials of subjects exhibiting decreased

gene expression, protein levels, or protein activity. Alternatively, the effectiveness of an agent, as determined by a screening assay, to decrease gene expression, protein levels or protein activity, can be monitored in clinical trials of subjects exhibiting increased gene expression, protein levels, or protein activity. In such clinical trials, expression or activity of a polypeptide of the invention and preferably, that of other polypeptide that have been
5 implicated in for example, a cellular proliferation disorder, can be used as a marker of the immune responsiveness of a particular cell.

For example, and not by way of limitation, genes, including those of the invention, that are modulated in cells by treatment with an agent (*e.g.*, compound, drug or small molecule) which modulates activity or expression of a polypeptide of the invention (*e.g.*, as
10 identified in a screening assay described herein) can be identified. Thus, to study the effect of agents on cellular proliferation disorders, for example, in a clinical trial, cells can be isolated and RNA prepared and analyzed for the levels of expression of a gene of the invention and other genes implicated in the disorder. The levels of gene expression (*i.e.*, a gene expression pattern) can be quantified by Northern blot analysis or RT-PCR, as
15 described herein, or alternatively by measuring the amount of protein produced, by one of the methods as described herein, or by measuring the levels of activity of a gene of the invention or other genes. In this way, the gene expression pattern can serve as a marker, indicative of the physiological response of the cells to the agent. Accordingly, this response state may be determined before, and at various points during, treatment of the individual
20 with the agent.

In a preferred embodiment, the present invention provides a method for monitoring the effectiveness of treatment of a subject with an agent (*e.g.*, an agonist, antagonist, peptidomimetic, protein, peptide, nucleic acid, small molecule, or other drug candidate identified by the screening assays described herein) comprising the steps of (i) obtaining a
25 pre-administration sample from a subject prior to administration of the agent; (ii) detecting the level of the polypeptide or nucleic acid of the invention in the preadministration sample; (iii) obtaining one or more post-administration samples from the subject; (iv) detecting the level the of the polypeptide or nucleic acid of the invention in the post-administration samples; (v) comparing the level of the polypeptide or nucleic acid of the invention in the
30 pre-administration sample with the level of the polypeptide or nucleic acid of the invention in the post-administration sample or samples; and (vi) altering the administration of the agent to the subject accordingly. For example, increased administration of the agent may be desirable to increase the expression or activity of the polypeptide to higher levels than detected, *i.e.*, to increase the effectiveness of the agent. Alternatively, decreased
35 administration of the agent may be desirable to decrease expression or activity of the polypeptide to lower levels than detected, *i.e.*, to decrease the effectiveness of the agent.

C. Methods of Treatment

The present invention provides for both prophylactic and therapeutic methods of treating a subject at risk of (or susceptible to) a disorder or having a disorder associated with aberrant expression or activity of a polypeptide of the invention, *e.g.*, cardiac infection (*e.g.*, myocarditis or dilated cardiomyopathy), central nervous system infection (*e.g.*, non-specific febrile illness or meningoencephalitis), pancreatic infection (*e.g.*, acute pancreatitis), respiratory infection (pneumonia), gastrointestinal infection, type I diabetes, cancer, familia hypercholesterolemia, treat hemophilia B, Marfan syndrome, protein S deficiency, allergy, inflammation, and gastroduodenal ulcer. Moreover, the polypeptides of the invention can be used to modulate cellular function, survival, morphology, proliferation and/or differentiation.

1. Prophylactic Methods

In one aspect, the invention provides a method for preventing in a subject, a disease or condition associated with an aberrant expression or activity of a polypeptide of the invention, by administering to the subject an agent which modulates expression or at least one activity of the polypeptide. Subjects at risk for a disease which is caused or contributed to by aberrant expression or activity of a polypeptide of the invention can be identified by, for example, any or a combination of diagnostic or prognostic assays as described herein. Administration of a prophylactic agent can occur prior to the manifestation of symptoms characteristic of the aberrancy, such that a disease or disorder is prevented or, alternatively, delayed in its progression. Depending on the type of aberrancy, for example, an agonist or antagonist agent can be used for treating the subject.

2. Therapeutic Methods

Another aspect of the invention pertains to methods of modulating expression or activity of a polypeptide of the invention for therapeutic purposes. The modulatory method of the invention involves contacting a cell with an agent that modulates one or more of the activities of the polypeptide. An agent that modulates activity can be an agent as described herein, such as a nucleic acid or a protein, a naturally-occurring cognate ligand of the polypeptide, a peptide, a peptidomimetic, or other small molecule. In one embodiment, the agent stimulates one or more of the biological activities of the polypeptide. Examples of such stimulatory agents include the active polypeptide of the invention and a nucleic acid molecule encoding the polypeptide of the invention that has been introduced into the cell. In another embodiment, the agent inhibits one or more of the biological activities of the polypeptide of the invention. Examples of such inhibitory agents include antisense nucleic acid molecules and antibodies. These modulatory methods can be performed *in vitro* (*e.g.*,

by culturing the cell with the agent) or, alternatively, *in vivo* (e.g., by administering the agent to a subject). As such, the present invention provides methods of treating an individual afflicted with a disease or disorder characterized by aberrant expression or activity of a polypeptide of the invention. In one embodiment, the method involves administering an agent (e.g., an agent identified by a screening assay described herein), or
5 combination of agents that modulates (e.g., upregulates or downregulates) expression or activity. In another embodiment, the method involves administering a polypeptide of the invention or a nucleic acid molecule of the invention as therapy to compensate for reduced or aberrant expression or activity of the polypeptide.

Stimulation of activity is desirable in situations in which activity or expression is
10 abnormally low or downregulated and/or in which increased activity is likely to have a beneficial effect. Conversely, inhibition of activity is desirable in situations in which activity or expression is abnormally high or upregulated and/or in which decreased activity is likely to have a beneficial effect.

The contents of all references, patents and published patent applications cited
15 throughout this application are hereby incorporated by reference.

Deposit of Clones

Clones containing cDNA molecules encoding human MANGO 003 were deposited with the American Type Culture Collection (ATCC® 10801 University Boulevard,
20 Manassas, VA 20110-2209) on March 30, 1999 as Accession Number 207178, as part of a composite deposit representing a mixture of three strains, each carrying one recombinant plasmid harboring a particular cDNA clone.

To distinguish the strains and isolate a strain harboring a particular cDNA clone, an aliquot of the mixture can be streaked out to single colonies on nutrient medium (e.g., LB
25 plates) supplemented with 100 g/ml ampicillin, single colonies grown, and then plasmid DNA extracted using a standard miniprep procedure. Next, a sample of the DNA miniprep can be digested with a combination of the restriction enzymes *Sal* I and *Not* I, and the resultant products resolved on a 0.8% agarose gel using standard DNA electrophoresis conditions. The digest liberates fragments as follows:

30 human MANGO 003 (clone EpthLa6a1): 3.2 kB

The identity of the strains can be inferred from the fragments liberated.

35 Clones containing cDNA molecules encoding human INTERCEPT 340, MANGO 347, and TANGO 272 were deposited with the American Type Culture Collection (ATCC®

10801 University Boulevard, Manassas, VA 20110-2209) on June 18, 1999 as Accession Number PTA-250, as part of a composite deposit representing a mixture of three strains, each carrying one recombinant plasmid harboring a particular cDNA clone.

To distinguish the strains and isolate a strain harboring a particular cDNA clone, an aliquot of the mixture can be streaked out to single colonies on nutrient medium (*e.g.*, LB plates) supplemented with 100 g/ml ampicillin, single colonies grown, and then plasmid DNA extracted using a standard miniprep procedure. Next, a sample of the DNA miniprep can be digested with a combination of the restriction enzymes *Sal* I and *Not* I, and the resultant products resolved on a 0.8% agarose gel using standard DNA electrophoresis conditions. The digest liberates fragments as follows:

human INTERCEPT 340 (clone EpI340): 3.3 kB
human MANGO 347 (clone EpM347): 1.4 kB
human TANGO 272 (clone EpT272): 5.0 kB

The identity of the strains can be inferred from the fragments liberated.

Clones containing cDNA molecules encoding human TANGO 295, TANGO 354, and TANGO 378 were deposited with the American Type Culture Collection (ATCC® 10801 University Boulevard, Manassas, VA 20110-2209) on June 18, 1999 as Accession Number PTA-249, as part of a composite deposit representing a mixture of three strains, each carrying one recombinant plasmid harboring a particular cDNA clone.

To distinguish the strains and isolate a strain harboring a particular cDNA clone, an aliquot of the mixture can be streaked out to single colonies on nutrient medium (*e.g.*, LB plates) supplemented with 100 g/ml ampicillin, single colonies grown, and then plasmid DNA extracted using a standard miniprep procedure. Next, a sample of the DNA miniprep can be digested with a combination of the restriction enzymes *Sal* I and *Not* I, and the resultant products resolved on a 0.8% agarose gel using standard DNA electrophoresis conditions. The digest liberates fragments as follows:

human TANGO 295 (clone EpT295): 1.5 kB
human TANGO 354 (clone EpT354): 1.8 kB
human TANGO 378 (clone EpT378): 3.3 kB

The identity of the strains can be inferred from the fragments liberated.

All publications, patents and patent applications mentioned in this specification are herein incorporated by reference into the specification to the same extent as if each individual publication, patent or patent application was specifically and individually indicated to be incorporated herein by reference.

5 Equivalents

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following Claims.

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25

30

35

MICROORGANISMS

Optional Sheet in connection with the microorganism referred to on pages ___, lines ___ of the description *

A. IDENTIFICATION OF DEPOSIT *

Further deposits are identified on an additional sheet *

Name of depositary institution *

American Type Culture Collection

Address of depositary institution (including postal code and country) *

10801 University Blvd.
Manassas, VA 20110-2209
USDate of deposit * March 30, 1999 Accession Number * 207178**B. ADDITIONAL INDICATIONS *** (leave blank if not applicable). This information is continued on a separate attached sheet**C. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE *** (if the indications are not all designated States)**D. SEPARATE FURNISHING OF INDICATIONS *** (leave blank if not applicable)

The indications listed below will be submitted to the International Bureau later * (Specify the general nature of the indications e.g., "Accession Number of Deposit")

E. ☒ This sheet was received with the International application when filed (to be checked by the receiving Office)
(Authorized Officer)☐ The date of receipt (from the applicant) by the International Bureau *

was

(Authorized Officer)

Form PCT/RO/134 (January 1981)

-116.2 -

International Application No: PCT/ /

Form PCT/RO/134 (cont.)

American Type Culture Collection

10801 University Blvd.
Manassas, VA 20110-2209
US

<u>Accession No.</u>	<u>Date of Deposit</u>
PTA-249	June 18, 1999
PTA-250	June 18, 1999

What is claimed is:

1. An isolated nucleic acid molecule selected from the group consisting of:
 - a) a nucleic acid molecule comprising a nucleotide sequence which is at least 55% identical to the nucleotide sequence of SEQ ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16,
5 18, 19, 21, 22, 24, 25, 27, 28, 30, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, or a complement thereof;
 - b) a nucleic acid molecule comprising a fragment of at least 300 nucleotides of
10 the nucleotide sequence of SEQ ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, 19, 21, 22, 24, 25, 27, 28, 30, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, or a complement thereof;
 - c) a nucleic acid molecule which encodes a polypeptide comprising the amino
15 acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence
20 encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250;
 - d) a nucleic acid molecule which encodes a fragment of a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC®
25 as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, wherein the fragment comprises at least 15 contiguous amino acids of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, 29, the amino acid sequence encoded
30 by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250; and
 - e) a nucleic acid molecule which encodes a naturally occurring allelic variant of
35 a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20,

23, 26, 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, wherein the nucleic acid molecule hybridizes to a
5 nucleic acid molecule comprising SEQ ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, 19, 21, 22, 24, 25, 27, 28, 30, or a complement thereof, under stringent conditions.

2. The isolated nucleic acid molecule of Claim 1, which is selected from the group consisting of:
10 a) a nucleic acid comprising the nucleotide sequence of SEQ ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, 19, 21, 22, 24, 25, 27, 28, 30, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, or a complement thereof; and
15 b) a nucleic acid molecule which encodes a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence
20 encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250.

3. The nucleic acid molecule of Claim 1 further comprising vector nucleic acid sequences.
25

4. The nucleic acid molecule of Claim 1 further comprising nucleic acid sequences encoding a heterologous polypeptide.

5. A host cell which contains the nucleic acid molecule of Claim 1.
30

6. The host cell of Claim 5 which is a mammalian host cell.

7. A non-human mammalian host cell containing the nucleic acid molecule of Claim 1.
35

8. An isolated polypeptide selected from the group consisting of:

- a) a fragment of a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29, wherein the fragment comprises at least 15 contiguous amino acids of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29;
- b) a naturally occurring allelic variant of a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29, the amino acid sequence
5 encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession
10 Number PTA-250, wherein the polypeptide is encoded by a nucleic acid molecule which hybridizes to a nucleic acid molecule comprising SEQ ID NOs: 1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, or a complement thereof under stringent conditions; and
- c) a polypeptide which is encoded by a nucleic acid molecule comprising a nucleotide sequence which is at least 55% identical to a nucleic acid comprising the nucleotide sequence of SEQ ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, 19, 21, 22, 24,
15 25, 27, 28, 30, or a complement thereof.
9. The isolated polypeptide of Claim 8 comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29.
10. The polypeptide of Claim 8 further comprising heterologous amino acid
20 sequences.
11. An antibody which selectively binds to a polypeptide of Claim 8.
12. A method for producing a polypeptide selected from the group consisting of:
25 a) a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession
30 Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250;
- b) a polypeptide comprising a fragment of the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as
35 Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, wherein the

fragment comprises at least 15 contiguous amino acids of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250; and

c) a naturally occurring allelic variant of a polypeptide comprising the amino acid sequence of SEQ ID NOs:2, 5, 8, 11, 14, 17, 20, 23, 26, or 29, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number 207178, the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-249, or the amino acid sequence encoded by the cDNA insert of the plasmid deposited with the ATCC® as Accession Number PTA-250, wherein the polypeptide is encoded by a nucleic acid molecule which hybridizes to a nucleic acid molecule comprising SEQ ID NOs:1, 3, 4, 6, 7, 9, 10, 12, 13, 15, 16, 18, 19, 21, 22, 24, 25, 27, 28, 30, or a complement thereof under stringent conditions;

comprising culturing the host cell of Claim 5 under conditions in which the nucleic acid molecule is expressed.

13. A method for detecting the presence of a polypeptide of Claim 8 in a sample, comprising:

- a) contacting the sample with a compound which selectively binds to a polypeptide of Claim 8; and
- b) determining whether the compound binds to the polypeptide in the sample.

14. The method of Claim 13, wherein the compound which binds to the polypeptide is an antibody.

15. A kit comprising a compound which selectively binds to a polypeptide of Claim 8 and instructions for use.

16. A method for detecting the presence of a nucleic acid molecule of Claim 1 in a sample, comprising the steps of:

- a) contacting the sample with a nucleic acid probe or primer which selectively hybridizes to the nucleic acid molecule; and
- b) determining whether the nucleic acid probe or primer binds to a nucleic acid molecule in the sample.

17. The method of Claim 16, wherein the sample comprises mRNA molecules and is contacted with a nucleic acid probe.

18. A kit comprising a compound which selectively hybridizes to a nucleic acid molecule of Claim 1 and instructions for use.

19. A method for identifying a compound which binds to a polypeptide of Claim 8 comprising the steps of:

a) contacting a polypeptide, or a cell expressing a polypeptide of Claim 8 with a test compound; and

b) determining whether the polypeptide binds to the test compound.

20. The method of Claim 19, wherein the binding of the test compound to the polypeptide is detected by a method selected from the group consisting of:

a) detection of binding by direct detecting of test compound/polypeptide binding;

b) detection of binding using a competition binding assay;

c) detection of binding using an assay for INTERCEPT 340-, MANGO 003-, MANGO 347-, TANGO 272-, TANGO 295-, TANGO 354-, or TANGO 378-mediated signal transduction.

21. A method for modulating the activity of a polypeptide of Claim 8 comprising contacting a polypeptide or a cell expressing a polypeptide of Claim 8 with a compound which binds to the polypeptide in a sufficient concentration to modulate the activity of the polypeptide.

22. A method for identifying a compound which modulates the activity of a polypeptide of Claim 8, comprising:

a) contacting a polypeptide of Claim 8 with a test compound; and

b) determining the effect of the test compound on the activity of the polypeptide to thereby identify a compound which modulates the activity of the polypeptide.

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Input file I340Athsa102b12; Output File I340Athsa102b12.pat
Sequence length 3284

GTGACCCACGCGTCCGTTATGTAACATACATTTTCCAGAAATTTAGTATATGATATGATTTTGTTCCTTCATC 79
CCTTTTCCAAGCAGTTTATTATGAAAATTTTCAAACATACAGCAATGTTGAGAAAATTTACAGTAAATGCCTATACC 158
CATTACCTAAATTTTACCATTAAATTTTACCCTGCTGGCATTATTGTGCTTATCCATCTACSTATCCCTCTCTCCCTT 237
CATTGGTGATTTTCTAAGTAAATGTAGGCCTCAGTACACTTCCTTCTGAATTCTTCAGCATGCACAACAGTATTATAT 316
TCCATTTTAAAAGAGCAATTCCTGATAGATTATATAGTTTGTAAAATGTTTCATATAGAGCTACAAAATTTATCTTT 395
TTGTTTCTTATTGTATGTCTAGGGTCTGAAGGGGATGCTGGCATTGTTGGGATATCAGGTCCTAAAGGTCTATTGGA 474
CACAGAGGAAACACTGGTCCCCTTGGCAGAGAAGGTATAATAGGCCAACAGGTAGAACTGGACCCAGAGGTGAAAAGG 553
GCTTTAGAGGTGAAACTGGTCTCAAGGACCAAGAGGTCAACCAGGGCTCCAGGTCCACCTGGAGCACCAGGCCCAAG 632
AAAGCAAATGGATATCAATGCTGCTATTCAAGCCTTGATTGAATCAAATACTGCCCTACAGATGGAGGTAACATATCTG 711
GTTTTATTATATTGGCACTGTCTCTCAATATACCAATTAACAGAGAAAATTTTGGAGGCCAAAATGTGACATTATC 790
TCAAAGATTGTATTTAAACAGATTGAAAATGTGAAACCATTCTCAAGAACAAAGTAAGTGATTTTGGTATAATTAAAC 869
AGAAATATATGCGTAGGATGTTTTGTAAGGAAAACATTTAAATCAAAAATTTAGTACTGTTATTTGTAAGGAATTTGGT 948
ACTATCCAAGAAAGTAGTTAAATGAGGTTAGCCATGTTTCTTAAATGAGATATATATATTATCACTACTCATTTATTT 1027
AAACTCTAATGATTCAATGTGTAATTTAAAAAACATAATACAGTAGACATAGCAATTCTTATGTTAGCTTGAAAACTAA 1106
ACTTGCAAATGTGAATTTAACCTCTTTAAAAGATTAAGGTTATTAAGCATACACATATGCCTATGCTTAAATATAAAC 1185

M E T H S S P A L A 10
TGTTCTTTACATTCTACTCACAACCTTACTACACATA ATG GAA ACA CAT TCT TCT CCT GCC TTG GCC 1251

H V G P Q D F F V Y I I L M M T W Q S Y 30
CAT GTT GGT CCT CAG GAT TTT TTT GTT TAT ATA ATT CTT ATG ATG ACT TGG CAG AGC TAC 1311

Q N T E V T L I D H S E E I F K T L N Y 50
CAG AAT ACT GAA GTG ACT TTA ATT GAC CAC AGT GAA GAG ATA TTC AAA ACC CTG AAC TAC 1371

FIG.1A

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L S N L L H S I K N P L G T R D N P A R	70
CTT AGC AAT TTA TTG CAC AGC ATC AAG AAT CCT CTT GGC ACA CGA GAT AAC CCA GCA CGA	1431
I C K D L L N C E Q K V S D G K Y W I D	90
ATC TGC AAA GAT TTA CTT AAC TGT GAA CAA AAA GTA TCA GAT GGA AAA TAC TGG ATT GAC	1491
P N L G C P S D A I E V F C N F S A G G	110
CCA AAT CTT GGC TGT CCT TCA GAT GCC ATT GAG GTT TTC TGC AAT TTC AGT GCT GGT GGC	1551
Q T C L P P V S V T K L E F G V G K V Q	130
CAG ACA TGC TTA CCT CCT GTT TCT GTA ACA AAG TTG GAG TTT GGA GTT GGG AAA GTC CAG	1611
M N F L H L L S S E A T H I I T I H C L	150
ATG AAC TTC CTT CAT TTA CTG AGT TCG GAA GCC ACC CAT ATC ATC ACC ATT CAC TGT CTA	1671
N T P R W T S T Q T S G P G L P I G F K	170
AAC ACC CCA AGG TGG ACA AGC ACA CAA ACA AGT GGC CCA GGA TTG CCT ATT GGT TTC AAG	1731
G W N G Q I F K V N T L L E P K V L S D	190
GGA TGG AAT GGC CAG ATT TTT AAA GTA AAC ACT CTA CTT GAA CCT AAA GTG CTT TCA GAT	1791
D C K I Q D G S W H K A T F L F H T Q E	210
GAC TGC AAG ATT CAA GAT GGC AGC TGG CAT AAG GCA ACA TTT CTT TTT CAC ACC CAG GAA	1851
P N Q L P V I E V Q K L P H L K T E R K	230
CCT AAT CAA CTT CCA GTG ATT GAA GTA CAA AAA CTT CCT CAT CTC AAA ACT GAA CGA AAG	1911
Y Y I D S S S V C F L *	242
TAT TAC ATT GAC AGC AGT TCT GTA TGC TTT CTG TAA	1947
AGTCTCTGAATTAGTTC CGAATTCAGGCTGTGGCCAGGTAATTGCTGCAGAGGGAGAAATAAGACAGACAGATACAGT	2026
CATTATGAAATGCATGTAATAAAGCATTGGCTAAATCTTAAAGAATCTCAGGAAGAACAGACTTCCTCCTAAGAAGGAG	2105
AAAAGGCATTTTTAAAGGACTATGATTGATAAAGTATTTAATTCTTTAAAAATTATATTCATCTCAGCTTTCTTAGAG	2184
AATTCCTAGAACTAAAAATTTATAAATATGGAATTCCTCAGGTATCTTATATTTTTGACTGAGTGGTAGTACCCAT	2263
TAGACAGCTGGAGATGCAGAGCACTATGGACCAATACTGGCTAATGCTTCCAGATGTGCACTGCTTCTGTCTAAAAATT	2342
ACAAGCCACAGTCTAATATGTCTTATTTTCCAAAACACTAAGCTGTATTTCAGGTCCCCGATGGGCATATACATCTTAGC	2421

FIG.1B

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CGGTGATACACTACCTCTTACGTGTGCCTCTTTGTGTGCTTGGTGCTCTTCGAAAACAAGGTGCTTATGGCTTTCA 2500
TAGACTATTTCCTTTTTCATCTTTGTCATTCTTTAAAAGTGTATGTACTGGTTACATCAAGATATGTTTGGTTGTTAG 2579
TACTTATTTTAATTTGTTTGGTCACACACTTAATAACACATGAAACTATTTATGTGAAGTCCTTGTTTTATTTAAAAT 2658
TCTCTTTGTGTATTTGGAATCAAAGCCAGCACATTGTAACCTGTGCTTGTACGCAAAAGAATTAGATTTCTTTGTTTTT 2737
GTTTTATTTTTAAATTGTTGTAAAAATTATTATAGGCCAGCTACATCTAGTAGTAGGTTTGGGTACAGATTGGGGT 2816
TGTGCCATACTGTTTTTAAAGTTCATGATCATCTGGAATGATACTTAGTGTATATATTTTGTAAAGTTTAAATTCAG 2895
CAAATTTTTTGAAATTGCTGCTGTTTTAAATTATAAAACCTTTATATTTCTGCTTTGTAGAAATTATATGTTTGTAGT 2974
ATTCATTGATTTCTTTCACTGTACTTAAATTTAGTGTTAGTACTTTAAAATTTTAAATTTACCAGTCTTTAAAGCAAC 3053
ATCCAGAAAAAAAAGTCTTTTCCCATTTAAATAGGCTCAGCCAGTTCATGTCGCCTTGTATCAGAGAAATATTA 3132
GTTCAATACTGAAAGAAAAATATTATACCTCTTGGTATCTAGAAAAGCTTGTTCATCCATTATAAATATATCTTTAGCC 3211
ACAGCAAACCACTTAACCTATCTATAATAAAAAATGTGCTTTAAATAAAAAAAAAAAAAAAAAAGGGCGGCCG 3284

FIG.1C

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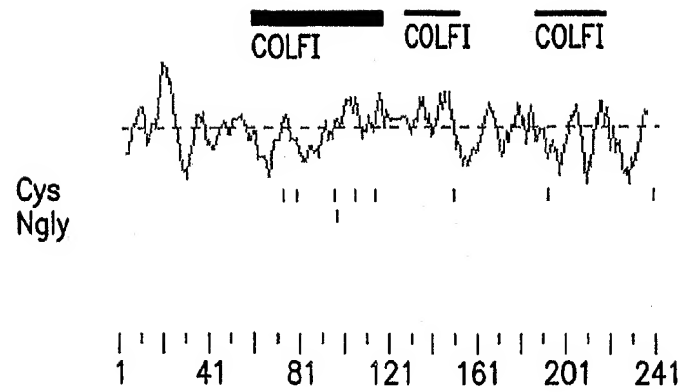


FIG.2

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COLFI: domain 1 of 3, from 58 to 116: score 110.3, E = 1.3e-42

```

      *->IksPeGksrknPARTCkDLfLchpefsGeYWiDPNqGCikDAikVf
      +k+P+G +r+nPAR CkDL c + ++G YWiDPN+GC+ DAi+Vf
INT340  58  IKNPLG-TRDNPARICKDLLNCEQKVSDGKYWIDPNLGCPSDAIEVF 103
      CnkrfetGvgeTCisp<-*
      Cn f +G g+TC +p
INT340 104 CN-FSAG-QQTCLPP 116
```

COLFI: domain 2 of 3, from 126 to 151: score 9.7, E = 0.0028

```

      *->isnvQITFLRLlSteAsQNITYhCKN<-*
      +++vQ+ FL LLS+eA iT hC N
INT340 126  VGKVQMNFLLHLLSSEATHIITIHLN 151
```

COLFI: domain 3 of 3, from 186 to 217: score 5.8, E = 0.09

```

      *->lvIGeDGCssrtgewgKTViEyeTkKtRLPIv<-*
      +vI D C+ g w K+ + + T+ + +LP +
INT340 186  KVL-SDDCKIQDGSWHKATFLFHTQEPNQLPVI 217
```

FIG.3

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Input file M003Athyo30d3; Output File M003AthYo30d3.pat

Sequence length 3169

	M T P S P	5
GTCGACCCACGCGTCCGCGCCCGCTGAGCCCCCGCCGAGGTCCGGACAGGCCGAG	ATG ACG CCG AGC CCC	71
L L L L L L P P L L L G A F P P A A A A		25
CTG TTG CTG CTC CTG CTG CCG CCG CTG CTG CTG GGG GCC TTC CCG CCG GCC GCC GCC GCC		311
R G P P K M A D K V V P R Q V A R L G R		45
CGA GGC CCC CCA AAG ATG GCG GAC AAG GTG GTC CCA CGG CAG GTG GCC CCG CTG GGC CGC		191
T V R L Q C P V E G D P P P L T M W T K		65
ACT GTG CCG CTG CAG TGC CCA GTG GAG GGG GAC CCG CCG CCG CTG ACC ATG TGG ACC AAG		251
D G R T I H S G W S R F R V L P Q G L K		85
GAT GGC CCG ACC ATC CAC AGC GGC TGG AGC CCG TTC CCG GTG CTG CCG CAG GGC CTG AAG		311
V K Q V E R E D A G V Y V C K A T N G F		105
GTG AAG CAG GTG GAG CCG GAG GAT GCC GGC GTG TAC GTG TGC AAG GCC ACC AAC GGC TTC		371
G S L S V N Y T L V V L D D I S P G K E		125
GGC AGC CTG AGC GTC AAC TAC ACC CTC GTC GTG CTG GAT GAC ATT AGC CCA GGC AAG GAG		431
S L G P D S S S G G Q E D P A S Q Q W A		145
AGC CTG GGC CCC GAC AGC TCC TCT GGG GGT CAA GAG GAC CCC GCC AGC CAG CAG TGG GCA		491
R P R F T Q P S K M R R R V I A R P V G		165
CGA CCG CCG TTC ACA CAG CCC TCC AAG ATG AGG CCG CCG GTG ATC GCA CCG CCC GTG GGT		551
S S V R L K C V A S G H P R P D I T W M		185
AGC TCC GTG CCG CTC AAG TGC GTG GCC AGC GGC CAC CCT CCG CCC GAC ATC ACG TGG ATG		611
K D D Q A L T R P E A A E P R K K K W T		205
AAG GAC GAC CAG GCC TTG ACG CCG CCA GAG GCC GCT GAG CCC AGG AAG AAG AAG TGG ACA		671
L S L K N L R P E D S G K Y T C R V S N		225
CTG AGC CTG AAG AAC CTG CCG CCG GAG GAC AGC GGC AAA TAC ACC TGC CCG GTG TCG AAC		731
R A G A I N A T Y K V D V I Q R T R S K		245
CGC GCG GGC GCC ATC AAC GCC ACC TAC AAG GTG GAT GTG ATC CAG CCG ACC CGT TCC AAG		791

FIG.4A

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P V L T G T H P V N T T V D F G G T T S	265
CCC GTG CTC ACA GGC ACG CAC CCC GTG AAC ACG ACG GTG GAC TTC GGG GGG ACC ACG TCC	851
F Q C K V R S D V K P V I Q W L K R V E	285
TTC CAG TGC AAG GTG CGC AGC GAC GTG AAG CCG GTG ATC CAG TGC CTG AAG CGC GTG GAG	911
Y G A E G R H N S T I D V G G Q K F V V	305
TAC GGC GCC GAG GGC CGC CAC AAC TCC ACC ATC GAT GTG GGC GGC CAG AAG TTT GTG GTG	971
L P T G D V W S R P D G S Y L N K L L I	325
CTG CCC ACG GGT GAC GTG TGG TCG CGG CCC GAC GGC TCC TAC CTC AAT AAG CTG CTC ATC	1031
T R A R Q D D A G M Y I C L G A N T M G	345
ACC CGT GCC CGC CAG GAC GAT GCG GGC ATG TAC ATC TGC CTT GGC GCC AAC ACC ATG GGC	1091
Y S F R S A F L T V L P D P K P P G P P	365
TAC AGC TTC CGC AGC GCC TTC CTC ACC GTG CTG CCA GAC CCA AAA CCG CCA GGG CCA CCT	1151
V A S S S S A T S L P W P V V I G I P A	385
GTG GCC TCC TCG TCC TCG GCC ACT AGC CTG CCG TGG CCG GTG GTC ATC GGC ATC CCA GCC	1211
G A V F I L G T L L L W L C Q A Q K K P	405
GGC GCT GTC TTC ATC CTG GGC ACC CTG CTC CTG TGG CTT TGC CAG GCC CAG AAG AAG CCG	1271
C T P A P A P P L P G H R P P G T A R D	425
TGC ACC CCC GCG CCT GCC CCT CCC CTG CCT GGG CAC CGC CCG CCG GGG ACG GCC CGC GAC	1331
R S G D K D L P S L A A L S A G P G V G	445
CGC AGC GGA GAC AAG GAC CTT CCC TCG TTG GCC GCC CTC AGC GCT GGC CCT GGT GTG GGG	1391
L C E E H G S P A A P Q H L L G P G P V	465
CTG TGT GAG GAG CAT GGG TCT CCG GCA GCC CCC CAG CAC TTA CTG GGC CCA GGC CCA GTT	1451
A G P K L Y P K L Y T D I H T H T H T H	485
GCT GGC CCT AAG TTG TAC CCC AAA CTC TAC ACA GAC ATC CAC ACA CAC ACA CAC ACA CAC	1511
S H T H S H V E G K V H Q H I H Y Q C *	505
TCT CAC ACA CAC TCA CAC GTG GAG GGC AAG GTC CAC CAG CAC ATC CAC TAT CAG TGC TAG	1571
ACGGCACCGTATCTGCAGTGGGCACGGGGGGCCGCCAGACAGGCAGACTGGGAGGATGGAGGACGGAGCTGCAGACG	1650

FIG.4B

AAGGCAGGGGACCCATGGCGAGGAGGAATGGCCAGCACCCCAGGCAGTCTGTGTGTGAGGCATAGCCCCCTGGACACACA 1729
 CACACAGACACACACTGCCTGGATGCATGTATGCACACACATGCGCGCACACGTGCTCCCTGAAGGCACACGTACGC 1808
 ACACACGCACATGCACAGATATGCCGCCTGGGCACACAGATAAGCTGCCCAAATGCACGCACACGCACAGAGACATGCC 1887
 AGAACATACAAGGACATGCTGCCTGAACATACACACGCACACCCATGCGCAGATGTGCTGCCTGGACACACACACAC 1966
 ACGGATATGCTGTCTGGACGCACACACGTGCAGATATGGTATCCGGACACACACGTGCACAGATATGCTGCCTGGACAC 2045
 ACAGATAATGCTGCCTTGACACACACATGCACGGATATTGCCTGGACACACACACACACGTGTGCACAGATATGCTG 2124
 TCTGGACACGCACACACATGCAGATATGCTGCCTGGACACACACTTCCAGACACACGTGCACAGGCGCAGATATGCTGC 2203
 CTGGACACACGCAGATATGCTGTCTAGTCACACACACACGCAGACATGCTGTCCGGACACACACACGCATGCACAGATA 2282
 TGCTGTCCGGACACACACACGCACGCAGATATGCTGCCTGGACACACACAGATAATGCTGCCTCAACACTCACACAC 2361
 GTGCAGATATTGCCTGGACACACACATGTGCACAGATATGCTGTCTGGACATGCACACACGTGCAGATATGCTGTCCGG 2440
 ATACACACGCACGCACACATGCAGATATGCTGCCTGGGCACACACTTCCGGACACACATGCACACACAGGTGCAGATAT 2519
 GCTGCCTGGACACACGCAGACTGACGTGCTTTTGGGAGGGTGTGCCGTGAAGCCTGCAGTACGTGTCCCGTGAGGCTCA 2598
 TAGTTGATGAGGACTTTCCCTGCTCCACCGTCACTCCCCCAACTGTGCCCGCCTCTGTCCCGCCTCAGTCCCCGCCT 2677
 CCATCCCCGCCTCTGTCCCTGGCCTTGGCGGCTATTTTTGCCACCTGCCTTGGGTGCCCAGGAGTCCCTACTGCTGT 2756
 GGGCTGGGGTTGGGGGCACAGCAGCCCCAAGCCTGAGAGGCTGGAGCCCATGGCTAGTGGCTCATCCCCACTGCATTCT 2835
 CCCCCTGACACAGAGAAGGGGCCTTGGTATTTATATTTAAGAAATGAAGATAATATTAATAATGATGGAAGGAAGACTG 2914
 GGTTCAGGGACTGTGGTCTCTCCTGGGGCCGGGACCCGCCTGGTCTTTTCAGCCATGCTGATGACCACACCCCGTCCA 2993
 GGCCAGACACCACCCCCACCCCACTGTCGTGGTGGCCCCAGATCTCTGTAATTTTATGTAGAGTTTGAGCTGAAGCCC 3072
 CGTATATTTAATTTATTTTGTAAACATGAAAGTGCAA 3151
 AAAAAAAGGGCGGCCGC 3169

FIG.4C

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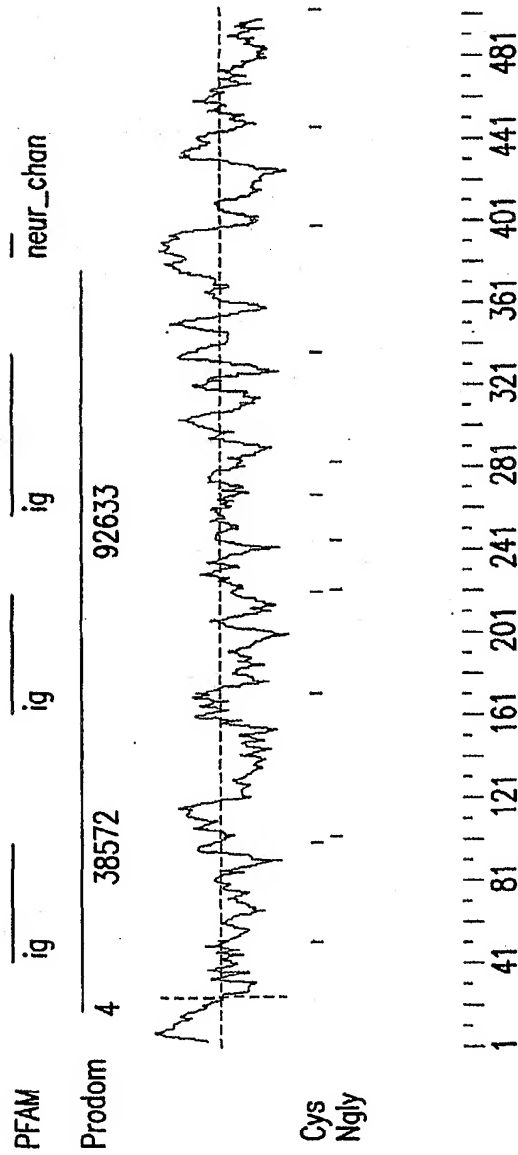


FIG.5

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ig: domain 1 of 3, from 44 to 101: score 36.4, E = 9.9e-10

```
*->GesvtLtCsvgfgpp.p.vtWlrngk.....lslti.s
G +v+L+C v g+p+p W+++g++ +++ ++ + + l ++
M003 44 GRTVRLQCPVE—GDPpPITMWTKDGRTihsgwsrfrvlpQGLKVkQ 88
```

```
vtpeDsgGtYtCvv<-*
v++eD+ G+Y C +
M003 89 VEREDA-GVYVCKA 101
```

ig: domain 2 of 3, from 165 to 223: score 48.9, E = 1.3e-13

```
*->GesvtLtCsvgfgpp.p.vtWlrngk.....lslti.
G+sv+L C +s g p+p+ttW +++t +++t +++++t +l ++
M003 165 GSSVRLKCVAS—GHPpPdITWMKDDQaltrpeaaepkrkkWTLSLk 209
```

```
svtpeDsgGtYtCvv<-*
+++peDs G YtC+v
M003 210 NLRPEDS-GKYTCRV 223
```

ig: domain 3 of 3, from 261 to 340: score 26.9, E = 8.8e-07

```
*->GesvtLtCsvgfgpp.p.vtWlrngk.....
G++ +++C vt ++ tp +HWl+ + + ++++++ + ++++
M003 261 GGTTSFQCKVR—SDVkpVlQWLKRVEygaegrhnstidvggqkfvv 305
```

```
.....lslti.svtpeDsgGtYtCvv<-*
++++ ++++++ l+i+++++D+ G Y C
M003 306 lptgdvwsrpdgsyINKLLlRARQDDA-GMYICLG 340
```

FIG.6

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neur_chan: domain 1 of 1, from 388 to 397: score 1.4, E = 9.7

->vfvlGTlgif<-
vf+IGTI ++
MO03 388 VFILGTLLW 397

FIG.7

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Input file M003jfmjf004c11; Output File M003jfmjf004c11.pat
Sequence length 1074

R V R P T G D V W S R P D G S Y L N K	19
CA CGC GTC CGG CCC ACG GGT GAT GTG TGG TCA CGG CCT GAT GGC TCC TAC CTC AAC AAG	59
L L I S R A R Q D D A G M Y I C L G A N	39
CTG CTC ATC TCT CGG GCC CGC CAG GAT GAT GCT GGC ATG TAC ATC TGC CTA GGT GCA AAT	119
T M G Y S F R S A F L T V L P D P K P P	59
ACC ATG GGC TAC AGT TTC CGT AGC GCC TTC CTC ACT GTA TTA CCA GAC CCC AAA CCT CCA	179
G P P M A S S S S S T S L P W P V V I G	79
GGG CCT CCT ATG GCT TCT TCA TCG TCA TCC ACA AGC CTG CCA TGG CCT GTG GTG ATC GCC	239
I P A G A V F I L G T V L L W L C Q T K	99
ATC CCA GCT GGT GCT GTC TTC ATC CTA GGC ACT GTG CTG CTC TGG CTT TGC CAG ACC AAG	299
K K P C A P A S T L P V P G H R P P G T	119
AAG AAG CCA TGT GCC CCA GCA TCT ACA CTT CCT GTG CCT GGG CAT CGT CCC CCA GGG ACA	359
S R E R S G D K D L P S L A V G I C E E	139
TCC CGA GAA CGC AGT GGT GAC AAG GAC CTG CCC TCA TTG GCT GTG GGC ATA TGT GAG GAG	419
H G S A M A P Q H I L A S G S T A G P K	159
CAT GGA TCC GCC ATG GCC CCC CAG CAC ATC CTG GCC TCT GGC TCA ACT GCT GGC CCC AAG	479
L Y P K L Y T D V H T H T H T H T C T H	179
CTG TAC CCC AAG CTA TAC ACA GAT GTG CAC ACA CAC ACA CAT ACA CAC ACC TGC ACT CAC	539
T L S C W R A R F I N T S M S T I S A K	199
ACG CTC TCA TGT TGG AGG GCA AGG TTC ATC AAC ACC AGC ATG TCC ACT ATC AGT GCT AAA	599
Y S E S P S T V S *	209
TAC AGC GAA TCT CCA AGC ACT GTG TCC TGA	629

FIG.8A

GGTAGGCATTGGGGCCAAGGCAACAGGTTGGGAGAATTGAGAACAATGGAGGAAGAGTATCTTAGGGTGCCTTATGG 708
TGGACACTCACAACTTGGCCATATAGATGTATGTACTACCAGATGAACAGCCAGCCAGATTACACACGCACATGTTT 787
AAACGTGTAAACGTGTGCACAACCTGCACACACAACCTGAGAAACCTTCAGGAGGATTTGTGGTGTGACTTTGCAGTGAC 866
ATGTAGCGATGGCTAGTTGAAGGAATCTCCCTCATGTCTTAGTGGTCATGGCCACTTCCCCACCCCTGCCCATCTGTGT 945
TCCTGCCTGGCCTTGGTGGTGCTTCCGTGTGCCCTGGGTTTTCCAGGAACCCCTATCAACCTGACTGGGGTGAGCAGTGC 1024
AGCCATGCNTGGAGGTTTGAGCCACCCTCCCCTTGCTAGAGAGAAGGCCN 1074

FIG.8B

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PFAM

Cys
Ngly



1 41 81 121 161 201

FIG.9

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Input file M347Alhbad295g12; Output File M347Alhbad295g12.pat
Sequence length 1423

	M P G P R V W G K Y L W	12
GTGACCCACGCGTCCGCCACGCGTCCGG	ATG CCT GGA CCC AGA GTG TGG GGG AAA TAT CTC TGG	66
R S P H S K G C P G A M W W L L L W G V		32
AGA AGC CCT CAC TCC AAA GGC TGT CCA GGC GCA ATG TGG TGG CTG CTT CTC TGG GGA GTC		126
L Q A C P T R G S V L L A Q E L P Q Q L		52
CTC CAG GCT TGC CCA ACC CGG GGC TCC GTC CTC TTG GCC CAA GAG CTA CCC CAG CAG CTG		186
T S P G Y P E P Y G K G Q E S S T D I K		72
ACA TCC CCC GGG TAC CCA GAG CCG TAT GGC AAA GGC CAA GAG AGC AGC ACG GAC ATC AAG		246
A P E G F A V R L V F Q D F D L E P S Q		92
GCT CCA GAG GGC TTT GCT GTG AGG CTC GTC TTC CAG GAC TTC GAC CTG GAG CCG TCC CAG		306
D C A G D S V T V S W G W G G S R Q D C		112
GAC TGT GCA GGG GAC TCT GTC ACA GTG AGC TGG GGA TGG GGG GGG TCC CGC CAG GAC TGT		366
G Q G D S R G C G K W R C P E S P I W R		132
GGC CAG GGA GAT TCC CGG GGT TGT GGG AAG TGG CGG TGC CCT GAA TCC CCC ATC TGG AGG		426
R D E F S M *		139
AGG GAT GAA TTT TCC ATG TAG		447
GGGCAGTCGGGCTTGGCTTACCGGGGAGCAGTGGTGGACCCAGGACACAGCCTCCCACCAGCGCCTCCGGGGCTGCCA		526
TCTGGGCCCCACAGAGCAAAGAGGGCAGCAAGCAGGCCCTGCGTTTGAAGGCTTATGAATGGACACACAAATCTTGCA		605
AATCTATGGAGCCAGGGCAGGGACGCACATATTGGTTGTTAAAAATATGTCATCATGTATTTGTTGAGTGCCTGCTCT		684
ATCAGGTGAGGAAGCTGGACACAAATAATAACAAAAGATTAAGTCACCGTTCACACTTACCTTGAAGAGCTATTACAA		763
AACTTCTAACGCCAAAGCCTTATTCAGAATAAGGACATTTTAAAAACAGTACTTGATGGAGTGATGCAAGCTTGCAGTC		842
CCAGCAGTATAGTCAGGAGACTGAGGCTGGAGGATCAGAGGGCTGGAGCCCAGGGTTCAAGGCCAGCCTAAGCAACATA		921

FIG.10A

GCAAGACCCCATCTCAAAAATAAGTAAATAATAAATAAAAAATAAAAGAGCACATTATCTTTTGATTTAAATTTTATTT 1000
ATATCAAAATGACATAAATTTTGAACTTTATTTTAAATTTTAAATTTTAAATTATTATGGATACATAATAGTTGTA 1079
AGACTTTTGTTTTTTAATTAAGTTTTCTAAGGCTGGGCGCAGTAGCTCATGTCTGTAGTCCCAGCACTTTGGGAGGC 1158
TGAGGCGAAAGAAGCACTTGAGCCCAGGAATTTGAGACCAGCCTGGGCAACATAGCAAGACCCCATCTCTACAAAAAAA 1237
TTTAAAAATTAGCCAAGTGTGGTGGCAGCACCTGTGGTCCCAGCTACAAGGACGCTGAAGTGAGAGGATCACTTGAG 1316
CCTGGAAGGTAGAGGCTGCAGTGAGCTCTGATCATGACACCGTACTCCAGCCTGGGTGACAGAGTGAGACCCCTGTCTCC 1395
AAAAAAAAAAAAAAAAAAGGGCGGCCGC 1423

FIG.10B

17/95

M347

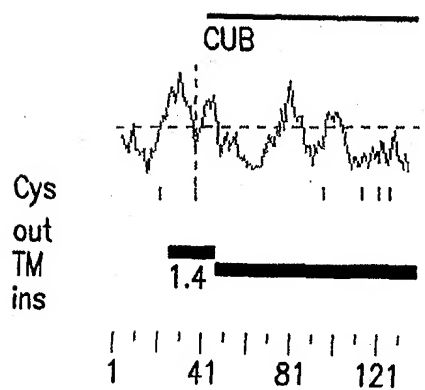


FIG. 11

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CUB: domain 1 of 1, from 40 to 136: score -17.7, E = 0.035

```

      *->CGgtldltessGsisSPnYPnrsdYppnkeCvWrlrappgyrvVeLt
      G +l+ +e + ++SP+YP+ +Y +e I ap+g+ V L
hm347 40 -GSVLLAQELPQQLTSPGYE--PYGKGQESSTDIKAPEGFA-VRLV 82

      FqdFdIEdhgapCryDyvEirDGdpss.plIG....rfCG....sgkPe
      FqdFdIE +++ C+ D+v + G ++s++ G+++r CG+ + ++P
hm347 83 FQDFDLEPSQD—CAGDSVTVSWGCGSrQDCGqgdsRCCGkwrcPESP- 129

      dirStsnrml i kFvsDasvskrGFkAty<-*
      + +D+ +
hm347 130-----IWRRDE-----F 136
```

FIG. 12

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Input file T272Athda89h3; Output File T272Athda89h3.pat
Sequence length 5036

GTGACCCACGCGTCCGCTCGAAGCGGGGACCTCGCCCCGCTCGGCTGTCCAGTCCTCCTCCTCGCAGACCCCGGC	79
GGTTCTACCCAGGCCCGCAGGGGAGACGGTGCCCCAAGGCAGGCTTCATATCCTGAACGCTGGGATCCCCAGGACAT	158
	M S 2
TCCCTGGCCCCCAGGCCCCAGGTCCCAGGCCCCAGGGCTGAGCTGTGGGCAGGCCCCACCTGGCCTCTGCA ATG TCA	235
P P L C P L L L L A V G L R L A G T L N	22
CCG CCT CTG TGT CCC CTC CTT CTC CTG GCT GTG GGC CTG CGG CTG GCT GGA ACT CTC AAC	295
P S D P N T C S F W E S F T T T T K E S	42
CCC AGT GAT CCC AAT ACC TGC AGC TTC TGG GAA AGC TTC ACT ACC ACC ACC AAG GAG TCC	355
H S R P F S L L P S E P C E R P W E G P	62
CAC TCC CGC CCC TTC AGC CTG CTC CCC TCA GAG CCC TGC GAG CGG CCC TGG GAG GGC CCC	415
H T C P S P Q T Q R K L L A S R D S F C	82
CAT ACT TGC CCC AGC CCA CAA ACT CAG AGG AAA CTC CTG GCT TCT AGG GAT TCA TTC TGC	475
M V C V G A G V Q W R D R S A L Q P Q T	102
ATG GTC TGT GTC GGG GCT GGA GTG CAG TGG CGA GAT CGT AGT GCA CTG CAA CCT CAA ACA	535
G N A L S M R P Q P R V L S G A P S L A	122
GGG AAT GCG CTT TCT ATG CGC CCT CAG CCC AGA GTG TTG AGT GGT GCC CCT TCC CTG GCC	595
S P G H T V V V K T D H R Q R L Q C C H	142
TCC CCT GGC CAC ACT GTG GTG GTG AAG ACG GAC CAC CGC CAG CGC CTG CAG TGC TGC CAT	655
G F Y E S R G F C V P L C A Q E C V H G	162
GGC TTC TAT GAG AGC AGG GGG TTC TGT GTC CCG CTC TGT GCC CAG GAG TGT GTC CAT GGC	715
R C V A P N Q C Q C V P G W R G D D C S	182
CGT TGT GTG GCA CCC AAT CAG TGC CAA TGT GTG CCA GGC TGG CGG GCC GAC GAC TGT TCC	775
S A P N C L Q P C T P G Y Y G P A C Q F	202
AGT GCC CCG AAC TGC CTT CAG CCC TGT ACC CCT GGC TAC TAT GGC CCT GCC TGC CAG TTC	835

FIG.13A

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R C Q C H G A P C D P Q T G A C F C P A	222
CGC TGC CAG TGC CAT GGG GCA CCC TGC GAT CCC CAG ACT GGA GCC TGC TTC TGC CCC GCA	895
E R T G P S C D V S C S Q G T S G F F C	242
GAG AGA ACT GGG CCC AGC TGT GAC GTG TCC TGT TCC CAG GGC ACT TCT GGC TTC TTC TGC	955
P S T H P C Q N G G V F Q T P Q G S C S	262
CCC AGC ACC CAT CCT TGC CAA AAT GGA GGT GTC TTC CAA ACC CCA CAG GGC TCC TGC AGC	1015
C P P G W M G T I C S L P C P E G F H G	282
TGC CCC CCT GGC TGG ATG GGC ACC ATC TGC TCC CTG CCC TGC CCA GAG GGC TTT CAC GGA	1075
P N C S Q E C R C H N G G L C D R F T G	302
CCC AAC TGC TCC CAG GAA TGT CGC TGC CAC AAC GGC GGC CTC TGT GAC CGA TTC ACT GGC	1135
Q C R C A P G Y T G D R C R E E C P V G	322
CAG TGC CGC TGC GCT CCG GGT TAC ACT GGG GAT CGG TGC CGG GAG GAG TGC CCG GTG GGC	1195
R F G Q D C A E T C D C A P D A R C F P	342
CGC TTT GGG CAG GAC TGT GCT GAG ACG TGC GAC TGC GCC CCG GAC GCC CGT TGC TTC CCG	1255
A N G A C L C E H G F T G D R C T D R L	362
GCC AAC GGC GCA TGT CTG TGC GAA CAC GGC TTC ACT GGG GAC CGC TGC ACG GAT CGC CTC	1315
C P D G F Y G L S C Q A P C T C D R E H	382
TGC CCC GAC GGC TTC TAC GGT CTC AGC TGC CAG GCC CCC TGC ACC TGC GAC CGG GAG CAC	1375
S L S C H P M N G E C S C L P G W A G L	402
AGC CTC AGC TGC CAC CCG ATG AAC GGG GAG TGC TCC TGC CTG CCG GGC TGG GCG GGC CTC	1435
H C N E S C P Q D T H G P G C Q E H C L	422
CAC TGC AAC GAG AGC TGC CCG CAG GAC ACG CAT GGG CCA GGG TGC CAG GAG CAC TGT CTC	1495
C L H G G V C Q A T S G L C Q C A P G Y	442
TGC CTG CAC GGT GGC GTC TGC CAG GCT ACC AGC GGC CTC TGT CAG TCC GCG CCG GGT TAC	1555
T G P H C A S L C P P D T Y G V N C S A	462
ACG GGC CCT CAC TGT GCT AGT CTT TGT CCT CCT GAC ACC TAC GGT GTC AAC TGT TCT GCA	1615

FIG.13B

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R	C	S	C	E	N	A	I	A	C	S	P	I	D	G	E	C	V	C	K	482
CGC	TGC	TCA	TGT	GAA	AAT	GCC	ATC	GCC	TGC	TCA	CCC	ATC	GAC	GGC	GAG	TGC	GTC	TGC	AAG	1675
E	G	W	Q	R	G	N	C	S	V	P	C	P	P	G	T	W	G	F	S	502
GAA	GGT	TGG	CAG	CGT	GGT	AAC	TGC	TCT	GTG	CCC	TGC	CCA	CCC	GGA	ACC	TGG	GGC	TTC	AGT	1735
C	N	A	S	C	Q	C	A	H	E	A	V	C	S	P	Q	T	G	A	C	522
TGC	AAT	GCC	AGC	TGC	CAG	TGT	GCC	CAT	GAG	GCA	GTC	TGC	AGC	CCC	CAA	ACT	GGA	GCC	TGT	1795
T	C	T	P	G	W	H	G	A	H	C	Q	L	P	C	P	K	G	Q	F	542
ACC	TGC	ACC	CCT	GGG	TGG	CAT	GGG	GCC	CAC	TGC	CAG	CTG	CCC	TGT	CCG	AAG	GGG	CAG	TTT	1855
G	E	G	C	A	S	R	C	D	C	D	H	S	D	G	C	D	P	V	H	562
GGA	GAA	GGT	TGT	CCC	AGT	CGC	TGT	GAC	TGT	GAC	CAC	TCT	GAT	GGC	TGT	GAC	CCT	GTT	CAT	1915
G	R	C	Q	C	Q	A	G	W	M	G	A	R	C	H	L	S	C	P	E	582
GGA	CGC	TGT	CAG	TGC	CAG	GCT	GGC	TGG	ATG	GGT	GCC	CGC	TGC	CAC	CTG	TCC	TGC	CCT	GAG	1975
G	L	W	G	V	N	C	S	N	T	C	T	C	K	N	G	G	T	C	L	602
GGC	TTA	TGG	GGA	GTC	AAC	TGT	ACC	AAC	ACC	TGC	ACC	TGC	AAG	AAT	GGG	GGC	ACC	TGT	CTC	2035
P	E	N	G	N	C	V	C	A	P	G	F	R	G	P	S	C	Q	R	S	622
CCT	GAG	AAT	GGC	AAC	TGC	GTG	TGT	GCA	CCC	GGA	TTT	CGG	GGC	CCC	TCC	TGC	CAG	AGA	TCC	2095
C	Q	P	G	R	Y	G	K	R	C	V	P	C	K	C	A	N	H	S	F	642
TGT	CAG	CCT	GGC	CGC	TAT	GGC	AAA	CGC	TGT	GTG	CCC	TGC	AAG	TGC	GCT	AAC	CAC	TCC	TTC	2155
C	H	P	S	N	G	T	C	Y	C	L	A	G	W	T	G	P	D	C	S	662
TGC	CAC	CCC	TGC	AAC	GGG	ACC	TGC	TAC	TGC	CTG	GCT	GGC	TGG	ACA	GGC	CCC	GAC	TGC	TCC	2215
Q	P	C	P	P	G	H	W	G	E	N	C	A	Q	T	C	Q	C	H	H	682
CAG	CCA	TGC	CCT	CCA	GGA	CAC	TGG	GGA	GAA	AAC	TGT	GCC	CAG	ACC	TGC	CAA	TGT	CAC	CAT	2275
G	G	T	C	H	P	Q	D	G	S	C	I	C	P	L	G	W	T	G	H	702
GGT	GGG	ACC	TGC	CAT	CCC	CAG	GAT	GGG	AGC	TGT	ATC	TGC	CCC	CTA	GGC	TGG	ACT	GGA	CAC	2335
H	C	L	E	G	C	P	L	G	T	F	G	A	N	C	S	Q	P	C	Q	722
CAC	TGC	TTA	GAA	GGC	TGC	CCT	CTG	GGG	ACA	TTT	GGT	GCT	AAC	TGC	TCC	CAG	CCA	TGC	CAG	2395

FIG.13C

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C G P G E K C H P E T G A C V C P P G H	742
TGT GGT CCT GGA GAA AAG TGC CAC CCA GAG ACT GGG GCC TGT GTA TGT CCC CCA GGG CAC	2455
S G A P C R I G I Q E P F T V M P T T P	762
AGT GGT GCA CCT TGC AGG ATT GGA ATC CAG GAG CCC TTT ACT GTG ATG CCG ACC ACT CCA	2515
V A Y N S L G A V I G I A V L G S L V V	782
GTA GCG TAT AAC TCG CTG GGT GCA GTG ATT GGC ATT GCA GTG CTG GGG TCC CTT GTG GTA	2575
A L V A L F I G Y R H W Q K G K E H H H	802
GCC CTG GTG GCA CTG TTC ATT GGC TAT CCG CAC TGG CAA AAA GGC AAG GAG CAC CAC CAC	2635
L A V A Y S S G R L D G S E Y V M P D V	822
CTG GCT GTG GCT TAC AGC AGC GGG CCG CTG GAC GGC TCC GAG TAT GTC ATG CCA GAT GTC	2695
P P S Y S H Y Y S N P S Y H T L S Q C S	842
CCT CCG AGC TAC AGT CAC TAC TAC TCC AAC CCC AGC TAC CAC ACC CTG TCG CAG TGC TCC	2755
P N P P P P N K V P G P L F A S L Q N P	862
CCA AAC CCC CCA CCC CCT AAC AAG GTT CCA GGC CCG CTC TTT GCC AGC CTG CAG AAC CCT	2815
E R P G G A Q G H D N H T T L P A D W K	882
GAG CCG CCA GGT GGG GCC CAA GGG CAT GAT AAC CAC ACC ACC CTG CCT GCT GAC TGG AAG	2875
H R R E P P P G P L D R G S S R L D R S	902
CAC CCG CCG GAG CCC CCT CCA GGG CCT CTG GAC AGG GGG AGC AGC CCG CTG GAC CGA AGC	2935
Y S Y S Y S N G P G P F Y D K G L I S E	922
TAC AGC TAT AGC TAC AGC AAT GGC CCA GGC CCA TTC TAC GAT AAA GGG CTC ATC TCT GAA	2995
E E L G A S V A S L S S E N P Y A T I R	942
GAG GAG CTC GGG GCC AGT GTG GCT TCC CTG AGC AGT GAG AAC CCA TAT GCC ACC ATC CCG	3055
D L P S L P G G P R E S S Y M E M K G P	962
GAC CTG CCC AGC TTG CCA GGG GGC CCC CCG GAG AGC AGC TAC ATG GAG ATG AAA GGC CCT	3115
P S G S A P R Q P P Q F W D S Q R R R Q	982
CCC TCA GGA TCT GCC CCC AGG CAG CCT CCT CAG TTT TGG GAC AGC CAG AGG CCG CCG CAA	3175

FIG.13D

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P Q P Q R D S G T Y E Q P S P L I H D R	1002
CCC CAG CCA CAG AGA GAC AGT GGC ACC TAC GAG CAG CCC AGC CCC CTG ATC CAT GAC CGA	3235
D S V G S Q P P L P P G L P P G H Y D S	1022
GAC TCT GTG GGC TCC CAG CCC CCT CTG CCT CCG GGC CTA CCC CCC GGC CAC TAT GAC TCA	3295
P K N S H I P G H Y D L P P V R H P P S	1042
CCC AAG AAC AGC CAC ATC CCT GGA CAT TAT GAC TTG CCT CCA GTA CGG CAT CCC CCA TCA	3355
P P L R R Q D R *	1051
CCT CCA CTT CGA CGC CAG GAC CGT TGA	3382
GGAGCCAGGATGGTATGGCAGAGGCCAGCACACCTGGCTGTTGCTGCTCAAGGCTGGGGACAGAGCCTAGTGTAACCCCT	3461
GCCAGGAGCAGGGAGTGGACCGGCAGGCTGTGAACATGAACAACGCTTAACAGAGCAAGTGATGGGAGCCTTGTTCTCTG	3540
GGTTCTACCATGGGAGACGCTGATCAGCAGGATGCCTGGCTCCCTTTCCCAACCCACTGCTCCCAAGGCCTCCAGGGCC	3619
CTGTGTACATAAACTGGTGGGTGGAAGTTGCTGGGTAAGTCTGATTTAGACATGGGTGTGGGTACCTTTTCTGTGC	3698
ATGCTCAGCCTGGGCTCTGTGGGTGTGTGTTTCTGTGATTTAGAGGGTACCAGGCAGGTTCTGTCTAGGGCACT	3777
TACCATTTAGTAGGGAGATGGAACCAACCAATTAAGTCTAGCAATAGCCTCCTAACTGGCCTCCTCCATTGATTCACT	3856
GAACCTTCCAATGCATGGCTCATAATTTCAAAATACAGGCTGGTTAGTTACTCCCTACCTGAAAGCCTTCATAGGTGCC	3935
TCTTTGCTCTTCTGCCAGTATCAAACTTTTGAAGGCCTTAAAGGCCCTGCTTTGCTGGCCCATCTGTCTCTCCAGCC	4014
TCACCTTGAAGTGTGTTCCGTGCACTGCACGCCAGTACACCGGCCTCTAGGTCTCTGTAGGCCACTCTTCTTTCTG	4093
GCACAGGGACCTGCACACCTGGAGTGGCTTCCCTCCCCCACTCGCCTGTTACCCCTGCTTTTCTTTACACCTCCTCC	4172
TCAGGGAAGTGGCCACCCCTCCGTACATCTTTCACAGCCCTGATTGCAGCTGTGTTCACTCACCAGGTACCTGCAGAAGG	4251
CCTACAGGGTGCCAGGCACTTCTTTAATGGGTCTTTCTTTATGTGATTATTTGATTAATCTCTGCCTCCCCCACTAGA	4330
CTGTAAGCTCCCTGAAGGCAAGAATCCTGTGCTTATGCTCAATATTAGCTCTCCCTTGGCACAGAGTAGGCACTCAACA	4409
AATGCTCCCCAAAAGGCTGAGTGGCTGACTGAATTAAGTACCAGTGACATGCAGTAAGTCTAAGATAGATGAGCCATC	4488

FIG.13E

TGTATGCTCTGACAGTTACAGACTGAATAAGTTGGAGACTTCCCTAAAGGGTGGCATTTCCTCAGGGTAACAACGCAGA 4567
GCTCAGGTGTGGGAAGGTGCCAGGGGCAGGGGTGCAGAGGGGCTGAGGCTGAGGGGGTGCAGAGGCTGGAGAAAGGAT 4646
AACAGGAGAGAGTATACAGGCATGCCTTGATTTATTGCACTTCACAGGTACAGAATTTTAAAGAAATTGAAGGTTTT 4725
GGGACATATATGTGACAGCAATAGGTTAAGAAAAGCAAAGCAGAGAAATTGAAGATTTGTGTCAACACTGCTTTAAGCA 4804
AATCTGTTGGCACCATTTTCCAATAGCATGTCCCATTTGGGTCTCTACATTGCATTTTGGTAATTGCTTGCAATAT 4883
TTCAAGCATTTTCATTGTTATTATATGTGTTATAGTGATCTGTGATCAGTGATCTTTGATATATTATTGTAATTGTTTC 4962
GGGGCGCCATGAACCGCACCCATATAACACGGTAAACTTAATCAGCAAAAAAAAAAAAAAAAAAAGGGCGGCCG 5036

FIG.13F

ECF-like

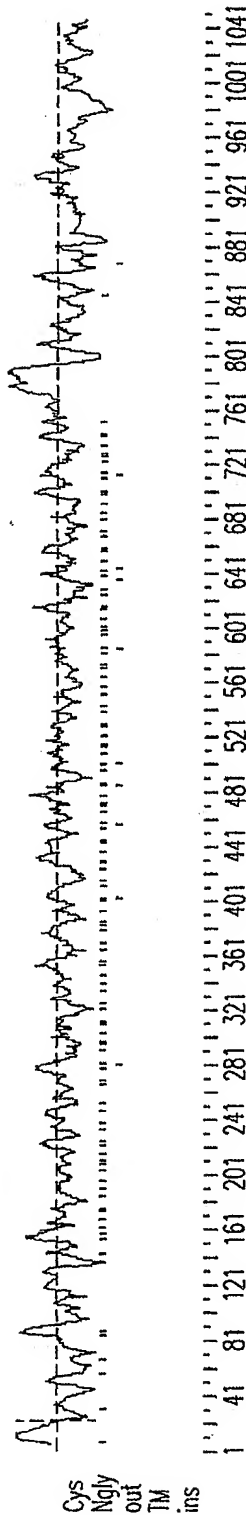


FIG. 14

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EGF: domain 1 of 14, from 151 to 181: score 14.0, E = 1.2

```

      *->Capnn..pCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC
      C p++ + C + G+Cv          +C+C pG      + G++C
hT272  151  CVPLCaqECVH-GRCVAPN-----QCQCVP-----WRGDDC 181

```

EGF: domain 2 of 14, from 200 to 229: score -2.2, E = 36

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
      C+ + C++ + C + g          C+Cp          tG+ C
hT272  200  CQFRCQCHG-APCDPQTG-----ACFCPAE-----RTGPSC 229

```

EGF: domain 3 of 14, from 242 to 272: score 16.0, E = 0.81

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
      C+++ pC+ngG+ + g          +C CppG      + G C
hT272  242  CPSTHPCQNGGVFTPTG-----SCSCPPG-----WMGTIC 272

```

EGF: domain 4 of 14, from 285 to 315: score 27.0, E = 0.00045

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
      C+++ C+ngG C g          +C+C+pG      ytG+rC
hT272  285  CSQECRCHNGGLCDRFTG-----QCRCAPG-----YTGDRC 315

```

FIG. 15A

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EGF: domain 5 of 14, from 328 to 358: score 18.0, E = 0.22

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyIGkrC<-
            Co+++ C   +++C +   g           C C +G   +tG+rC
hT272  328  CAETCDCAFDARCFPANG-----ACLCEHG-----FTGDRC   358

```

EGF: domain 6 of 14, from 378 to 404: score 7.4, E = 4.9

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyIGkrC<-
            C+ +   +   C++ g           +C C pG   ++G +C
hT272  378  CDRE----HSLSCHPMNG-----ECSCLPG-----WAGLHC   404

```

EGF: domain 7 of 14, from 417 to 447: score 29.2, E = 9.3e-05

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyIGkrC<-
            C++++ C++gG+C+ t g           C+C+pG   yIG++C
hT272  417  CQEHCLCLHGGVCQATSG-----LCQCAPG-----YTGPLHC   447

```

EGF: domain 8 of 14, from 460 to 490: score 6.0, E = 6.5

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyIGkrC<-
            C+ + C n   C + g           +C+C+HG   ++ +C
hT272  460  CSARCSCEAIACSPIDG-----ECVCKEG-----WQRCNC   490

```

FIG.15B

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EGF: domain 9 of 14, from 503 to 533: score 15.9, E = 0.82

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
            C+ + C + ++C + g          C+C+pG      ++G +C
hT272  503  CNASCQCAHEAVCSPQTG-----ACTCTPG-----WHGAHC  533

```

DSL: domain 1 of 1, from 518 to 576: score -20.5, E = 6.8

```

      *->WstdkhiggrtsIGfnleyrirvtCdenYYGegCnkFCrPrdDafgH
            +t + + + +      + +      C + +GegC+ C+      H
hT272  518  -QTGACTCTPG-----WHGAHCQLPCPKGQFEGGCASRCDGD-----H 554

```

```

            yt.Cd.enGnkICleGWkGeyC<-*
            + +Cd+ +G+ +C +GW+G C
hT272  555  SDgCDpVHGRCQCQAGWMGARC  576

```

EGF: domain 10 of 14, from 546 to 576: score 11.7, E = 2

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
            Ca+ + C++ C +++g          +C+C+ G      + G rC
hT272  546  CASRCDGDHSDGCDPVHG-----RCQCQAG-----WMGARC  576

```

EGF: domain 11 of 14, from 589 to 619: score 17.9, E = 0.24

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
            C+ ++ C+ngGtC++ g          C+C+pG      + G+ C
hT272  589  CSNTCTCKNGGTCLPENG-----NCVCAPG-----FRGPSC  619

```

FIG.15C

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EGF: domain 12 of 14, from 632 to 661: score 18.0, E = 0.23

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyGkrC<-
            C p  C n+ +C+++ g          tC C G      +tG++C
hT272  632  CVPC-KCANHSFCHPSNG-----TCYCLAG-----WTGPDC  661

```

EGF: domain 13 of 14, from 674 to 704: score 27.1, E = 0.00042

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyGkrC<-
            C+++ C++gGtC++ g          +C+Cp G      +tG++C
hT272  674  CAQTCQCHHGGTCHPQDG-----SCICPLG-----WTGHHC  704

```

EGF: domain 14 of 14, from 717 to 747: score 1.7, E = 16

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyGkrC<-
            C+++ C g +C++ g          C+CppG      +G C
hT272  717  CSQPCQCGPGCKHPETG-----ACVCPG-----HSGAPC  747

```

FIG.15D

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Input file t272Atmzb62c4; Output File t272Atmzb62c4.pat
Sequence length 2569

S T H A S G D P V H G Q C R C Q A G W	19
G TCG ACC CAC GCG TCC GGT GAC CCT GTT CAT GGA CAG TGC CGA TGT CAG GCT GGT TGG	58
M G T R C H L P C P E G F W G A N C S N	39
ATG GGC ACA CGC TGC CAC CTG CCT TGC CCG GAG GGC TTT TGG GGA GCC AAC TGC AGT AAC	118
T C T C K N G G T C V S E N G N C V C A	59
ACC TGT ACC TGC AAG AAT GGT GGT ACC TGT GTG TCT GAG AAT GGC AAC TGC GTG TGC GCA	178
P G F R G P S C Q R P C P P G R Y G K R	79
CCA GGG TTC CGA GGC CCC TCC TGC CAG AGG CCC TGC CCG CCT GGT CGC TAT GGC AAA CGC	238
C V Q C K C N N N H S S C H P S D G T C	99
TGT GTG CAA TGC AAG TGT AAC AAC AAC CAT TCT TCC TGC CAC CCA TCG GAC GGG ACC TGC	298
S C L A G W T G P D C S E A C P P G H W	119
TCC TGC CTG GCG GGC TGG ACA GGC CCT GAC TGC TCC GAG GCA TGT CCC CCA GGC CAC TGG	358
G L K C S Q L C Q C H H G G T C H P Q D	139
GGA CTC AAA TGC TCC CAA CTC TGC CAG TGT CAT CAT GGT GGG ACC TGC CAC CCC CAG GAT	418
G S C I C T P G W T G P N C L E G C P P	159
GGG AGC TGT ATC TGC ACG CCA GGC TGG ACT GGA CCC AAC TGC TTG GAA GGC TGC CCA CCA	478
R M F G V N C S Q L C Q C D L G E M C H	179
AGA ATG TTT GGT GTC AAC TGC TCC CAG CTA TGT CAG TGT GAT CTC GGA GAG ATG TGC CAC	538
P E T G A C V C P P G H S G A D C K M G	199
CCA GAG ACT GGG GCT TGT GTC TGT CCC CCA GGA CAC AGT GGT GCA GAC TGC AAA ATG GGA	598
S Q E S F T I M P T S P V T H N S L G A	219
AGC CAG GAG TCC TTC ACC ATA ATG CCC ACC TCT CCC GTG ACC CAT AAC TCA CTG GGT GCA	658
V I G I A V L G T L V V A L I A L F I G	239
GTG ATT GGC ATT GCA GTA CTG GGA ACC CTC GTG GTG GCC CTG ATA GCA CTG TTC ATT GGC	718

FIG. 16A

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Y R Q W Q K G K E H E H L A V A Y S T G	259
TAC CGC CAG TGG CAA AAG GGC AAG GAA CAT GAG CAC TTG GCA GTG GCT TAC AGC ACT GGG	778
R L D G S D Y V M P D V S P S Y S H Y Y	279
CGG CTG GAT GGC TCT GAT TAC GTC ATG CCA GAT GTC TCT CCG AGC TAT AGT CAC TAC TAC	838
S N P S Y H T L S Q C S P N P P P P N K	299
TCC AAC CCC AGC TAC CAC ACA CTG TCT CAG TGT TCT CCT AAC CCC CCG CCC CCT AAC AAG	898
V P G S Q L F V S S Q A P E R P S R A H	319
GTC CCA GGC AGT CAG CTC TTT GTC AGC TCT CAG GCC CCT GAG CCG CCA AGC AGA GCC CAC	958
G R E N H T T L P A D W K H R R E P H D	339
GGG CGT GAG AAC CAT ACC ACA CTG CCC GCT GAC TGG AAG CAC CCG CCG GAG CCC CAT GAC	1018
R G A S H L D R S Y S C S Y S H R N G P	359
AGA GGC GCC AGC CAC CTG GAC CGA AGC TAT AGC TGT AGC TAT AGC CAC AGG AAT GGC CCA	1078
G P F C H K G P I S E E G L G A S V M S	379
GGA CCA TTC TGT CAT AAA GGT CCC ATC TCT GAA GAG GGA CTA GGG GCA AGC GTT ATG TCC	1138
L S S E N P Y A T I R D L P S L P G E P	399
CTG AGC AGT GAG AAC CCC TAT GCT ACC ATC CGA GAC CTG CCC AGC CTG CCT GGG GAA CCC	1198
R E S G Y V E M K G P P S V S P P R Q S	419
CGA GAA AGT GGC TAT GTG GAG ATG AAA GGA CCT CCA TCA GTG TCC CCT CCC AGG CAG TCT	1258
L H L R D R Q Q R Q L Q P Q R D S G T Y	439
CTT CAT CTC CCG GAC AGG CAG CAG CCG CAA CTG CAG CCA CAG AGG GAC AGC GGC ACC TAT	1318
E Q P S P L S H N E E S L G S T P P L P	459
GAG CAG CCC AGC CCC TTG AGC CAT AAT GAA GAG TCT TTG GGC TCC ACG CCC CCG CTT CCT	1378
P G L P P G H Y D S P K N S H I P G H Y	479
CCA GGC CTG CCT CCT GGT CAC TAC GAC TCC CCC AAG AAC AGC CAT ATC CCT GGA CAC TAT	1438
D L P P V R H P P S P P S R R Q D R *	498
GAC TTG CCT CCA GTA CCG CAT CCT CCA TCC CCT CCA TCC CCG CCG CAG GAC CCG TGA	1495

FIG.16B

AGAGCCGGCATGGTATGGGAGCGTGCCTATGTACCTTGCCAGGAGCAGGGACTGGACCAGCAGGCCACGAACAGAAACA 1574
CTTGGTGAAGTGAACAGAGACGGACTGTGGCCCTGTGCTTCCACCGAGGGAGACACTAGTTGACAAAGTGTCTAACCCCT 1653
CTTTTCCAACCCACTGCTCAAGTCCCTGTGGACATAAGCTGGTGGGCAGAATGTTGTTGTACAAGTGTGATTTTAGATC 1732
GATTTTTTTTTAAAGTATGTGTGGGTACCTTTTCTGTGTGTATGCTCAGGCAGGCTGTGTGTCTCTAGTTGGCTTT 1811
AGAGGGAGTCAGGTATAGTTCTGCCTTCTGCACCTTCCATCTTATCTAGTAGTCAGCTTCCAAGCTTAACTAGTTAGA 1890
GCTCCACCAGCAGCAGGCCCTAACTACCTGCCTGCCCTTCACCCAGTAATCCTCCATGTCTTTGCTCAGAGGATTGCTC 1969
CCCGACTCTGGTGTGTCTCCTCGGTACGCCTTGACGGTCTGCAGTCTCCCTTTCCCGCTTGCTTCATTCTTTCCCA 2048
GAATGAAGGCTGTCTGCCACCCTACTTCCCAGCCCAGGAATTGGCACATCTAAGTTCAGCCTTCCTAAGTTACCCGTTG 2127
AGTCCTGCTTGCCTTCACATATTCACAGAACACCCACCCACATCTGCTTCATAGCTACTCTCTTCTCCACGTACCC 2206
ACAGAAGGCAGAAGTGGTACCAGGCAAGAAGATGGGATTGTTGCATTTTGTGTTTTGAGACTCTGTCTCACTATG 2285
TAGTCCTGGCTGGCCTGGAAGTCAAGAGCTCTGCCTGCCTCTGCCTCTTGAGTGTGGGTTAACGGCTCAGGGTCACA 2364
TGCACAGCTCAAGCTGCACTCCGATGTGCTTTCCCTGTTGCTAGATTAGCGTCTGCCTCCCCCTAGTGGAGAGGCTGA 2443
TCGCCAGCTCTCTGATGCAGGACTCTGGTGTGTTAGGCTCACTCACTATTGGTTTCCTTGGCACAGGGTAGTCACTCAAT 2522
AAATGTTCTCTAAAAGCTGAAAAAAAAAAAAAAAAAGGGCGGCCCG 2569

FIG.16C

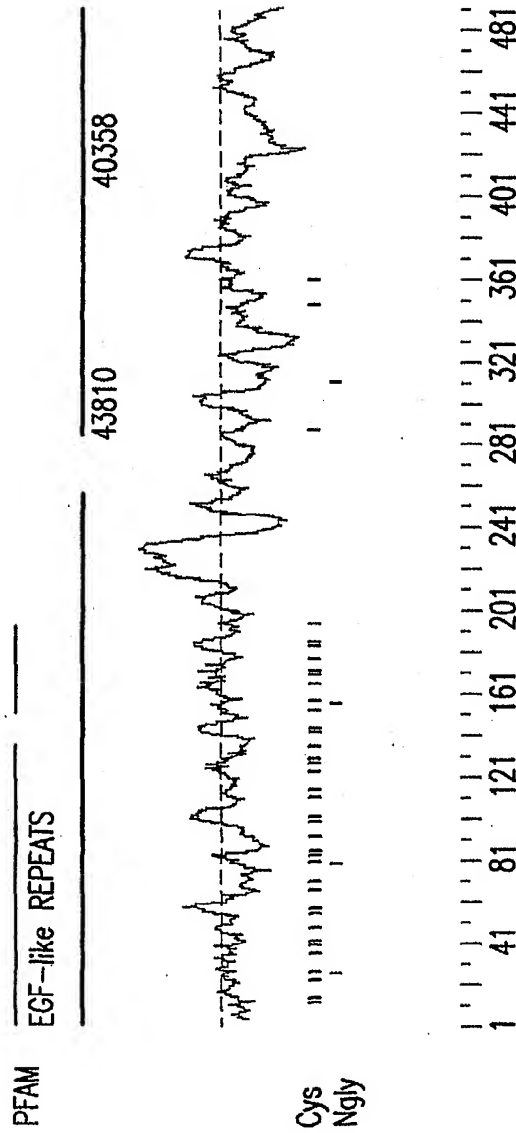


FIG.17

Input file T295Athyb23d9; Output File T295Athyb23d9.pat
Sequence length 1497

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GTGACCCACGCGTCCGGCTCCCAGCCCACCCCAACAGACACAGCGTAGCCCGGGCCAGCTCTTAAGGAGTTCAGGA 79
GTGAGAAGAGGCCCTCAGAGATCTGACAGCCTAGGAGTGGGTGGACACCACCTCAGCCCACTGAGCAGGAGTCACAGCA 158
CGAAGACCAAGCGCAAAGCGACCCCTGCCCTCCATCCTGACTGCTCCTCCTAAGAGAG ATG GCA CCG GCC AGA 231
M A P A R 5
A G F C P L L L L L L L G L W V A E I P 25
GCA GGA TTC TGC CCC CTT CTG CTG CTT CTG CTG CTG GGG CTG TGG GTG GCA GAG ATC CCA 291
V S A K P K G M T S S Q W F K I Q H M Q 45
GTC AGT GCC AAG CCC AAG GGC ATG ACC TCA TCA CAG TGG TTT AAA ATT CAG CAC ATG CAG 351
P S P Q A C N S A M K N I N K H T K R C 65
CCC AGC CCT CAA GCA TGC AAC TCA GCC ATG AAA AAC ATT AAC AAG CAC ACA AAA CGG TGC 411
K D L N T F L H E P F S S V A A T C Q T 85
AAA GAC CTC AAC ACC TTC CTG CAC GAG CCT TTC TCC AGT GTG GCC GCC ACC TGC CAG ACC 471
P K I A C K N G D K N C H Q S H G P V S 105
CCC AAA ATA GCC TGC AAG AAT GGC GAT AAA AAC TGC CAC CAG AGC CAC GGG CCC GTG TCC 531
L T M C K L T S G K Y P N C R Y K E K R 125
CTG ACC ATG TGT AAG CTC ACC TCA GGG AAG TAT CCG AAC TGC AGG TAC AAA GAG AAG CGA 591
Q N K S Y V V A C K P P Q K K D S Q Q F 145
CAG AAC AAG TCT TAC GTA GTG GCC TGT AAG CCT CCC CAG AAA AAG GAC TCT CAG CAA TTC 651
H L V P V H L D R V L * 157
CAC CTG GTT CCT GTA CAC TTG GAC AGA GTC CTT TAG 687

```

FIG.18A

GTTTCCAGACTGGCTTGCTCTTTGGCTGACCTTCAATTCCCTCTCCAGGACTCCGCACCACTCCCCTACACCCAGAGCA 766
TTCTCTTCCCCTCATCTCTGGGGCTGTTCTGGTTACGCTCTGCTGGGAGGCTGAAGCTGACACTCTGGTGAGCTGA 845
GCTCTAGAGGGATGGCTTTTCATCTTTTTGTTGCTGTTTTCCAGATGCTTATCCCCAAGAAACAGCAAGCTCAGGTCT 924
GTGGGTCCCTGGTCTATGCCATTGCACATGTCTCCCCTGCCCCCTGGCATTAGGGCAGCATGACAAGGAGAGGAAATA 1003
AATGGAAAGGGGCATATGGGATTTGTGGACACAGCTGTTTCTGTTCTGAACTAGAAGTCTTCCCAGCTCTGACGTG 1082
GCAGTGAGGTGACCTGAAGGAAAGAAAAATATAAATAAATACCACTTCATATTTGTATAGAATCCTCTAATCCCTTGTC 1161
ACATAGACTTGACAGGGATTGTATGCCTTCTTTATGGATGAGGAAATTAAGGTTTTAGAAAGCTTAATGAATTAAAGAG 1240
CTTGCTAATTAGTTAGTAGCAGAACCTGGACTTGAACCTAGGTCTCCTTGCTCTAAATACAGTGACCTTCTACTCTA 1319
CCAGTTGCGCAAGAAAGAAGTCACTGTTACAGAGGCAAGCGGTGAAGTGAAGAGTCACTCATGAAGAAACGAGTG 1398
CTCTGAAGAGCCAGTTACCCTGTGTTGGCTGCAATAAAGGTATTACCTCTCTAGCCAAAAAAAAAAAAAAAAAAAAAA 1477
AAAAAAAAAAAAAAAAAAAAA 1497

FIG.18B

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T295

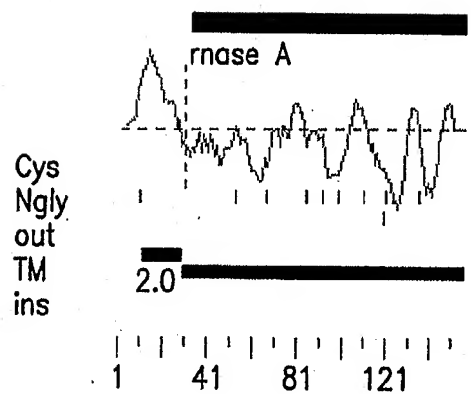


FIG.19

*->qesrAqkFlrQHIDspktsssnpnYCNqMMdkrRnmtqgrCKpvNTF
 + ++ q+F++QH+ ++s + CN +M k++n rCK+ NTF
 32 GMTSSQWFKIQHM---QPSPQA---CNSAM-KNINKHTKRCKDLNTF 71

vHesladVkaVCsqkNvtCKNGqkNCyqSkssfqiTdCr1tggsqkyPnC
 +He++++V a C ++ + CKNG kNC+qS+ ++++T C+lt+g yPnC
 72 LHEPFSSVAATCQTPKIACKNGDKNCHQSHGPVSLTMCKLTSGK--YPNC 119

rYrtsastkhIiVACEgrd.rddPyynPyvPVHFDasv<.*
 rY+ + ++k ++VAC +++++d+ ++ vPVH+D++
 120 RYKEKRQNKSYVVACKPPQkKDSQQFH-LVPVHLDRVL 156

FIG.20

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Input file T354Athla42a4; Output File T354Athla42a4.pat
Sequence length 1788

M P L L																				4	
GTCGACCCACGGCGTCCGGCCAGGCTCCACTGAGGGGAACGGGGACCTGTCTGAAGAGAAG																				ATG CCC CTG CTG	73
T L Y L L L F W L S G Y S I A T Q I T G																				24	
ACA CTC TAC CTG CTC CTC TTC TGG CTC TCA GGC TAC TCC ATT GCC ACT CAA ATC ACC GGT																				133	
P T T V N G L E R G S L T V Q C V Y R S																				44	
CCA ACA ACA GTG AAT GGC TTG GAG CGG GGC TCC TTG ACC GTG CAG TGT GTT TAC AGA TCA																				193	
G W E T Y L K W W C R G A I W R D C K I																				64	
GGC TGG GAG ACC TAC TTG AAG TGG TGG TGT CGA GGA GCT ATT TGG CGT GAC TGC AAG ATC																				253	
L V K T S G S E Q E V K R D R V S I K D																				84	
CTT GTT AAA ACC AGT GGG TCA GAG CAG GAG GTG AAG AGG GAC CGG GTG TCC ATC AAG GAC																				313	
N Q K N R T F T V T M E D L M K T D A D																				104	
AAT CAG AAA AAC CGC ACG TTC ACT GTG ACC ATG GAG GAT CTC ATG AAA ACT GAT GCT GAC																				373	
T Y W C G I E K T G N D L G V T V Q V T																				124	
ACT TAC TGG TGT GGA ATT GAG AAA ACT GGA AAT GAC CTT GGG GTC ACA GTT CAA GTG ACC																				433	
I D P A S T P A P T T P T S T T F T A P																				144	
ATT GAC CCA GCG TCG ACT CCT GCC CCC ACC ACG CCT ACT TCC ACT ACG TTT ACA GCA CCA																				493	
V T Q E E T S S S P T L T G H H L D N R																				164	
GTC ACC CAA GAA GAA ACT AGC AGC TCC CCA ACT CTG ACC GGC CAC CAC TTG GAC AAC AGG																				553	
H K L L K L S V L L P L I F T I L L L L																				184	
CAC AAG CTC CTG AAG CTC AGT GTC CTC CTG CCC CTC ATC TTC ACC ATA TTG CTG CTG CTT																				613	
L V A A S L L A W R M M K Y Q Q K A A G																				204	
TTG GTG GCC GCC TCA CTC TTG GCT TGG AGG ATG ATG AAG TAC CAG CAG AAA GCA GCC GGG																				673	
M S P E Q V L Q P L E G D L C Y A D L T																				224	
ATG TCC CCA GAG CAG GTA CTG CAG CCC CTG GAG GGC GAC CTC TGC TAT GCA GAC CTG ACC																				733	

FIG. 21A

L Q L A G T S P R K A T T K L S S A Q V	244
CTG CAG CTG GCC GGA ACC TCC CCG CGA AAG GCT ACC ACG AAG CTT TCC TCT GCC CAG GTT	793
D Q V E V E Y V T M A S L P K E D I S Y	264
GAC CAG GTG GAA GTG GAA TAT GTC ACC ATG GCT TCC TTG CCG AAG GAG GAC ATT TCC TAT	853
A S L T L G A E D Q E P T Y C N M G H L	284
GCA TCT CTG ACC TTG GGT GCT GAG GAT CAG GAA CCG ACC TAC TGC AAC ATG GGC CAC CTC	913
S S H L P G R G P E E P T E Y S T I S R	304
AGT AGC CAC CTC CCC GGC AGG GGC CCT GAG GAG CCC ACG GAA TAC AGC ACC ATC AGC AGG	973
P *	306
CCT TAG	979
CCTGCACTCCAGGCTCCTTCTTGGACCCAGGCTGTGAGCACACTCCTGCCTCATCGACCGTCTGCCCCCTGCTCCCCCT	1058
CATCAGGACCAACCCGGGGACTGGTGCTCTGCCTGATCAGCCAGCATTGCCCTAGCTCTGGGTTGGGCTTGGGGCCA	1137
AGTCTCAGGGGGCTTCTAGGAGTTGGGGTTTTCTAAACGTCCCTCCTCTCTACATAGTTGAGGAGGGGGCTAGGGAT	1216
ATGCTCTGGGGCTTTCATGGGAATGATGAAGATGATAATGAGAAAAATGTTATCATTATTATCATGAAGTACCATTATC	1295
ATAATACAATGAACCTTTATTTATTGCCTACCACATGTTATGGGCTGAATAATGGCCCCAAAGATATCTGTGTCCTAA	1374
TCCTCAGAACTTGTGACTGTTACCTTCTGTGGCAGAAAGGACAGTGCAGATGTATGTAAGTTAAGGACTTTGAGATAG	1453
AGAGGTTATTCTTGCTGATTACAGTGGGCCCAAAATATCACCACAAGGTCCTCATAAGAAAGAGGCCAGAAGGTCAA	1532
GAGGTAGAGACAAAGTGATGATGGAAGTGGACGTGGGTGTGACGTGAGCAGGGGCCATGAATGCCCGAGCCTTCAGATG	1611
CCAGAAAGGGAAGGAATGGATTCCCCTGCCTGGAGCCTCCAAAAGAAACCAGCCCTGCCACGCCTTGACTTGAGCCC	1690
ATTGAACTGATCTTGAGCTCCTGGCCTCCAGAATTGCAGGAGAATAAATTTGTGTTGTTTTAAAAAAAAAAAAAAAA	1769
AAAAAGGGCGGCCCTAGA	1788

FIG.21B

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T354

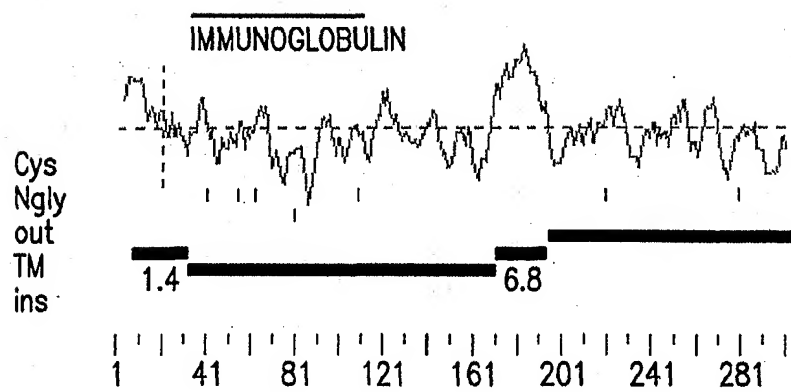


FIG.22

```

*->GesvtLtCsvsgfgppgsvtWf.....kngk.lgpsllgysyr1
    ++s+t +C ++ + + +++ W+ ++ ++ k l ++ s +
33  RGS LTVQCVYR--SGWETYLKWWCrgaiwRDCKiLVK--TSGSEQEV 75

esgekanlsegrfsis.....sltLtissvekeDsGtYtCvv<.*
    ++          r+si +++++++t+t+ ++ k D+ tY+C
76 KRD-----RVS IKdnqknrTFTVTMEDLMKTDADTYWCGI    110

```

FIG.23

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Input file T378Athta28f4; Output File T378Athta28f4.pat
Sequence length 3258

	M D H C G A L F L	9
CACGCGTCCGCCAGTTCTTGGAGGAGACTCTGCACAGGCC	ATG GAT CAC TGT GGT GCC CTT TTC CTG	68
C L C L L T L Q N A T T E T W E E L L S		29
TGC CTG TGC CTT CTG ACT TTG CAG AAT GCA ACA ACA GAG ACA TGG GAA GAA CTC CTG AGC		128
Y M E N M Q V S R G R S S V F S S R Q L		49
TAC ATG GAG AAT ATG CAG GTG TCC AGG GGC CGG AGC TCA GTT TTT TCC TCT CGT CAA CTC		188
H Q L E Q M L L N T S F P G Y N L T L Q		69
CAC CAG CTG GAG CAG ATG CTA CTG AAC ACC AGC TTC CCA GGC TAC AAC CTG ACC TTG CAG		248
T P T I Q S L A F K L S C D F S G L S L		89
ACA CCC ACC ATC CAG TCT CTG GCC TTC AAG CTG AGC TGT GAC TTC TCT GGC CTC TCG CTG		308
T S A T L K R V P Q A G G Q H A R G Q H		109
ACC AGT GCC ACT CTG AAG CGG GTG CCC CAG GCA GGA GGT CAG CAT GCC CGG GGT CAG CAC		368
A M Q F P A E L T R D A C K T R P R E L		129
GCC ATG CAG TTC CCC GCC GAG CTG ACC CGG GAC GCC TGC AAG ACC CGC CCC AGG GAG CTG		428
R L I C I Y F S N T H F F K D E N N S S		149
CGG CTC ATC TGT ATC TAC TTC TCC AAC ACC CAC TTT TTC AAG GAT GAA AAC AAC TCA TCT		488
L L N N Y V L G A Q L S H G H V N N L R		169
CTG CTG AAT AAC TAC GTC CTG GGG GCC CAG CTG AGT CAT GGG CAC GTG AAC AAC CTC AGG		548
D P V N I S F W H N Q S L E G Y T L T C		189
GAT CCT GTG AAC ATC AGC TTC TGG CAC AAC CAA AGC CTG GAA GGC TAC ACC CTG ACC TGT		608
V F W K E G A R K Q P W G G W S P E G C		209
GTC TTC TGG AAG GAG GGA GCC AGG AAA CAG CCC TGG GGG GGC TGG AGC CCT GAG GGC TGT		668
R T E Q P S H S Q V L C R C N H L T Y F		229
CGT ACA GAG CAG CCC TCC CAC TCT CAG GTG CTC TGC CGC TGC AAC CAC CTC ACC TAC TTT		728

FIG.24A

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A V L M Q L S P A L V P A E L L A P L T	249
GCT GTT CTC ATG CAA CTC TCC CCA GCC CTG GTC CCT GCA GAG TTG CTG GCA CCT CTT ACG	788
Y I S L V G C S I S I V A S L I T V L L	269
TAC ATC TCC CTC GTG GGC TGC AGC ATC TCC ATC GTG GCC TCG CTG ATC ACA GTC CTG CTG	848
H F H F R K Q S D S L T R I H M N L H A	289
CAC TTC CAT TTC AGG AAG CAG AGT GAC TCC TTA ACA CGC ATC CAC ATG AAC CTG CAT GCC	908
S V L L L N I A F L L S P A F A M S P V	309
TCC GTG CTG CTC CTG AAC ATC GCC TTC CTG CTG AGC CCC GCA TTC GCA ATG TCT CCT GTG	968
P G S A C T A L A A A L H Y A L L S C L	329
CCC GGG TCA GCA TGC ACG GCT CTG GCC GCT GCC CTG CAC TAC GCG CTG CTC AGC TGC CTC	1028
T W M A I E G F N L Y L L L G R V Y N I	349
ACC TGG ATG GCC ATC GAG GGC TTC AAC CTC TAC CTC CTC CTC GGG CGT GTC TAC AAC ATC	1088
Y I R R Y V F K L G V L G W G A P A L L	369
TAC ATC CGC AGA TAT GTG TTC AAG CTT GGT GTG CTA GGC TGG GGG GCC CCA GCC CTC CTG	1148
V L L S L S V K S S V Y G P C T I P V F	389
GTG CTG CTT TCC CTC TCT GTC AAG AGC TCG GTA TAC GGA CCC TGC ACA ATC CCC GTC TTC	1208
D S W E N G T G F Q N M S I C W V R S P	409
GAC AGC TGG GAG AAT GGC ACA GGC TTC CAG AAC ATG TCC ATA TGC TGG GTG CGG AGC CCC	1268
V V H S V L V M G Y G G L T S L F N L V	429
GTG GTG CAC AGT GTC CTG GTC ATG GGC TAC GGC GGC CTC ACG TCC CTC TTC AAC CTG GTG	1328
V L A W A L W T L R R L R E R A D A P S	449
GTG CTG GCC TGG GCG CTG TGG ACC CTG GCG AGG CTG CGG GAG CGG GCG GAT GCA CCA AGT	1388
V R A C H D T V T V L G L T V L L G T T	469
GTC AGG GCC TGC CAT GAC ACT GTC ACT GTG CTG GGC CTC ACC GTG CTG CTG GGA ACC ACC	1448
W A L A F F S F G V F L L P Q L F L F T	489
TGG GCC TTG GCC TTC TTT TCT TTT GGC GTC TTC CTG CTG CCC CAG CTG TTC CTC TTC ACC	1508

FIG.24B

I L N S L Y G F F L F L W F C S Q R C R	509
ATC TTA AAC TCG CTC TAC GGT TTC TTC CTT TTC CTG TGG TTC TGC TCC CAG CGG TGC CGC	1568
S E A E A K A Q I E A F S S S Q T T Q *	529
TCA GAA GCA GAG GCC AAG GCA CAG ATA GAG GCC TTC AGC TCC TCC CAA ACA ACA CAG TAG	1628
TCGGGGCCTCCTGGCCTGGAATCCTCAGCCTCTCTGGCCGCCAGTAGCCTGAGGCTACGGCTCCTGCTAGAGAGGGTGG	1707
CAGGCCTGCTGCTGGACCCAGAGGCCACTGTGACCGCCAAGGGGCCTTTTCCACTTCCACGGCCTCTCCAGGCACTGA	1786
GGGGAAGGCATTGCTCTACCTCTCCCTGACATTTTGCTCCGGGGCAGATCCAACCTTACCTGGGGCAGCAAACCTTTGTC	1865
CTGGTACCTGGGCCCAGCTCGCCAGGGATGTGGGCAGAGCACCAGCCTGGGCATCAGGAAGCCAAGTTTCAAGGACTGT	1944
CTTTGAGTCTGTCTGTATGACCTTGGGCCTGCCACTTCTCACAGACCCTAGGTATCCACAGCTGTGACATGGGGCAAG	2023
CGGCTTTGTTTCAGCCTAACCCAGGAGCTTAGTAAAAATTGCATAAGACCAGGGGAAGAGTGTACGCGTGGGGTGGGA	2102
ATTCCCGCGGCCTCCACCTGCTTGCTAGGGGCAGGATCTCATTAGGCTGCCCTGGAAGCACCTGCTTGGCCCTGCCAC	2181
CTTCCTCCAGGGGAGGGCCAGATGGCATCCTGGCTTGGGGCGGTGGGACCTACCCAGGCTCTGAGACTTTACTGGCCT	2260
ATGCCTGAGGCCTCTTTTCTTTAACTCCCTAAATTATGATGACTCCAAGTCCAAGCCACCTTCCCAAAGATTGGGA	2339
GGTTCCGCGTTCCAGAGGCTCCTCTGCGGTGCTCCCAAGACTTCCATAGACCATCTGGACCAGTAGCCCATCCCGC	2418
AGTTTTCTTGGGGCAGAGGAAACGCTTCTTTCTCCTCCAGCTGAATCAGCTGGATCCCAGTGTCTGGCTGTTTGGT	2497
GATTGGGCAAGATTGAATTTGCCAGGTAGGCGTGAGAGTGTGGGTTTTAAATTGGAAGCTCAGGCCATAGTTTCAGAG	2576
AATCACCTTACCCAGACCTTCATGAGACAGTGCTCATGAAGCCAGTGGTTTCCAGAACGAACACTAGGGCGCACC	2655
GTTGGTCCACACTCAGAGGCCCTTGGCGCCAAGACTGCATCTAGAATCGCTCAAACACCTGTTTGAGACCCCATGCAC	2734
CAGCTGGAGGGGCCGTAAGTGCAGGACTGCGCCTACTGAGTGACCCATTTCTCCAGGAGGAAAGGCAAGACACGCTTA	2813
CACGGCCATTTGTCTCTTTTCCCAATGCGGCGGTGCACTTTCGCTCTTGGGGCTGCACCCAGACATAGCTGGCACCA	2892

FIG.24C

GAGCAGGGTGCTCAGGTGGTGGGTGCTCAGGGCCCTGCCCCAGGCCACTGGGCCGTTTTGATGACCTCGAAGGTCACAG 2971
GCAGAAAATAGGAGCAGGATTTCCCCTGGGAAAAGTTCTCCTGGGACATCTTCTGCTCTTCTGTACATTTCTAGATGC 3050
AAATAACTCCTTCACCAGGCAGTGAGTGGCGTAGGCTCTGGAGCCAGGCTGCCTGGGCTCCAATGCCAGCTCTGCCACT 3129
TGCTAGCTGTGAGACTGTGGACAAACCACTCAGCCTCTGTGTGCCTCAGTTTTCTATTTGTAAAATAGAGGCCATAGT 3208
GGTACCTATTTTGAAGACTAAGTAAAAGAATTCAAATAAAGAGACTTGGC 3258

FIG.24D

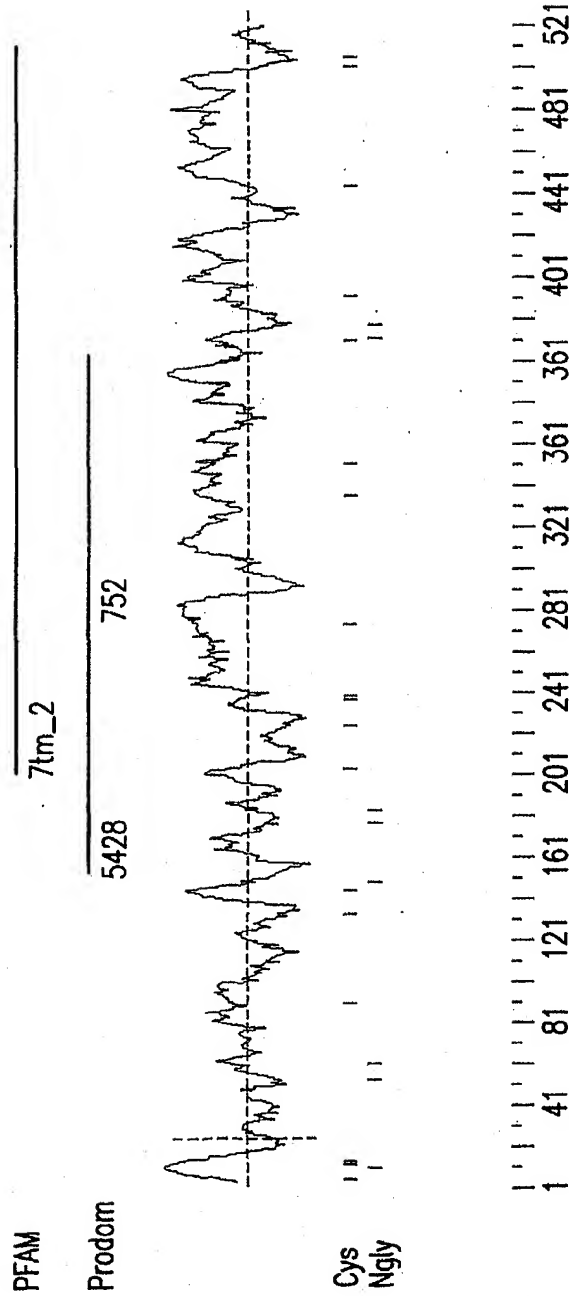


FIG.25

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```

*->CnrtWDgitC..Wpdt....ppGelVvvpCPkyfygfsdqttdtgn
      +tC W+ + +++p+G ++ C      + +q  + +
187 -----LTCvfWKEGarkqPWGGWSPEGC-----RTEQ---PSH 216

      vsRnCtedGsWsepppsNrtWrnysaCgeddpeesekkkkyylvikiY
      ++ C+ +   +++      + ++ +   +++ + 1 +1
217 SQVLCRCNH--LTYFA-----VLMQLSPALVPAELLAPLTYIS 252

      tvGYSISLaaLlvAvvILl1FRKLhtlwpdnadgalevgapWGAPfqvrr

      +vG S+S++a 1+ v++   FRk      + +
253 LVGCSISIVASLITVLLHFHFRKQS-----DSL----- 280

      SirCtRnyIHmNLFISFILrAasvfikdavlksevssdeperLssrcsls
      tR IHmNL +S +L +++ ++ a   s v+ ++
281 ----TR--IHMNLHASVLLLNIAFLLSPAFAMSPVPGSA----- 313

      tgqvvvvgCk11vvfQfqYcvmtNffWlLvEGlYlhtLLvvttffsErkylw
      C +1 ++ ++Y++++ +W+ +EG L+ LL  +   ++y +
314 -----CTALAAA-LHYALLSCLTWMaIEGFNLYLLLGRVY---NIYIR 352

      wYl....1IGWGvPlVfvvWaivRl1fedtgCWdsnGLAmFPEAKmCiW
      Y+ + +++GWG+P++ v      v++ ++ +C++++ F
353 RYVfk1gVLGWGAPALLVLLSLSVKSSVY-GPCTIPV----FDSWENG TG 397

      msdnshlwWIikgPiLlsilV.....NfflFinIirILvtKLraa
      n+++ W+ + P++ s+1V + ++ ++ N++++ ++ L + LR+
398 F-QNMSICWV-RSPVVHSLVvmgyggltslfNLVVLAWALWTL-RRLRER 444

      qtgetdqrqYsqYrkLaKSTL1LIPLfGIhyvvFafrPsndarGvlrkik
      + +      + +   L L L+G++ + +f+++ v+ +
445 ADAPSVR-----ACHDTVTVLGLTVLLGTTWALAFFSFG-----VFLLPQ 484

      lyfelSLgSFQGFfVAv1YCF1NgEVQaEirrrW<.*
      1++ L+S+ Gff ++ F+ + ++E +
485 LFLFTILNSLYGFF--LFLWFCSQRCRSEAEAKA      516

```

FIG.26

[illegible]

FIG. 27A

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```

      710      720      730      740      750      760      770
inputs ACAAGGTGGATGTGATCCAGCGGACCCGTTCCAAGCCCGTGCTCACAGGCACGCACCCCGTGAACACGAC
.....

      780      790      800      810      820      830      840
inputs GGTGGACTTCGGGGGGACACGTCTTCCAGTGCAAGGTGCGCAGCGACGTGAAGCCGGTGATCCAGTGG
.....

      850      860      870      880      890      900      910
inputs CTGAAGCGCGTGGAGTACGGCGCCGAGGGCCGCCACAACCTCCACCATCGATGTGGGCGGCCAGAAAGTTTG
.....

      920      930      940      950      960      970      980
inputs TGGTGCTGCCCACGGGTGACGTGTGGTGC GCGCCGACGGCTCCTACCTCAATAAGCTGCTCATCACCCG
.....GCCCACGGGTGATGTGTGGTACGCGCCTGATGGCTCCTACCTCAACAAGCTGCTCATCTCTCG
      20      30      40      50      60      70

      990      1000      1010      1020      1030      1040      1050
inputs TGCCCGCCAGGACGATGCGGGCATGTACATCTGCCTTGGCGCCAACACCATGGGCTACAGCTTCCGCAGC
.....GGCCCGCCAGGATGATGCTGGCATGTACATCTGCCTAGGTGCAAATACCATGGGCTACAGTTTCCGTAGC
      80      90      100      110      120      130      140

      1060      1070      1080      1090      1100      1110      1120
inputs GCCTTCCTCACCGTGCTGCCAGACCCAAAACCGCCAGGGCCACCTGTGGCCTCCTCGTCTCGGCCACTA
.....GCCTTCCTCACTGTATTACCAGACCCCAAACCTCCAGGGCCTCCTATGGCTTCTTCATCGTCATCCACAA
      150      160      170      180      190      200      210

      1130      1140      1150      1160      1170      1180      1190
inputs GCCTGCCGTGGCCCGTGGTGCATCGGCATCCAGCCGCGCTGTCTTCATCCTGGGCACCCCTGCTCCTGTG
.....GCCTGCCATGGCCTGTGGTGATCGGCATCCAGCTGGTGTCTTTCATCCTAGGCATGTGCTGCTCTG
      220      230      240      250      260      270      280

      1200      1210      1220      1230      1240      1250      1260
inputs GCTTTGCCAGGCCCAGAAGAAGCCGTGCACCCCCGCGCCTGCCCCTCCCCTGCCTGGGCACCGCCGCGCG
.....GCTTTGCCAGACCAAGAAGAAGCCATGTGCCCCAGCATCTACACTTCTGTGCCTGGGCATCGTCCCCCA
      290      300      310      320      330      340      350

      1270      1280      1290      1300      1310      1320      1330
inputs GGGACGGCCCGGACCGCAGCGGAGACAAGGACCTTCCCTCGTTGGCGCCCTCAGCGCTGGCCCTGGTG
.....GGGACATCCCGAGAACGCAGTGGTGACAAGGACCTGCCCTCATTGGC-----TG
      360      370      380      390      400

      1340      1350      1360      1370      1380      1390      1400
inputs TGGGGCTGTGTGAGGAGCATGGGTCTCCGGCAGCCCCCAGCACTTACTGGGCCCAGGCCAGTTGCTGG
.....TGGGCATATGTGAGGAGCATGGATCCGCCATGGCCCCCAGCACATCTGGCCTCTGGCTCAACTGCTGG
      410      420      430      440      450      460      470

```

FIG.27B

```

      1410      1420      1430      1440      1450      1460
inputs CCCTAAGTTGTACCCCAAACCTCTACACAGACATCCACACACACACA--CACACAC--TCTCACACACA
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
      CCCCAGCTGTACCCCAAAGCTATACACAGATGTGCACACACACACACATACACACACCTGCACTCACACG
      480      490      500      510      520      530      540

      1470      1480      1490      1500      1510
inputs CTCACACGT-GGAGGGCAAGGT-C-----CACCAGCACATCCACTATCAGTGC-----
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
      CTCTCATGTTGGAGGGCAAGGTTTCATCAACACCAGCATGTCCACTATCAGTGCTAAATACAGCGAATCTC
      550      560      570      580      590      600      610

inputs -----
      CAAGCACTGTGTCC

```

FIG.27C

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```
970      980      990      1000      1010      1020      1030
GTGCTGCCACGGGTGACGTGTGGTCGCGGCCGGACGGCTCTACCTCAATAAGCTGCTCATCACCCCTG
:::
GTCCGGCCACGGGTGATGTGTGGTCACGGCCTGATGGCTCTACCTCAACAAGCTGCTCATCTCTCGGG
10      20      30      40      50      60      70

1040      1050      1060      1070      1080      1090      1100
CCCCCAGGACGATGCGGGCATGTACATCTGCCTTGGCGCCAACACCATGGGCTACAGCTTCCGCAGCGA
:::
CCCCCAGGATGATGCTGGCATGTACATCTGCCTAGGTGCAAATACCATGGGCTACAGTTTCCGTAGCGC
80      90      100      110      120      130      140

1110      1120      1130      1140      1150      1160      1170
CTTCCTCACCGTGCTGCCAGACCCAAAACCGCCAGGGCCACCTGTGGCCTCCTCGTCTCGGCCACTAGC
:::
CTTCCTCACTGTATTACCAGACCCCAAACCTCCAGGGCCTCCTATGGCTTCTTCATCGTCATCCACAAGC
150      160      170      180      190      200      210

1180      1190      1200      1210      1220      1230      1240
CTGCCGTGGCCCGTGGTCATCGGCATCCAGCCGGCGCTGTCTTCATCCTGGGCACCCCTGCTCCTGTGGC
:::
CTGCCATGGCCTGTGGTGATCGGCATCCAGCTGGTGCTGTCTTCATCCTAGGCACTGTGCTGCTCTGGC
220      230      240      250      260      270      280

1250      1260      1270      1280      1290      1300      1310
TTTGCCAGGCCCAGAAGAAGCGTGACCCCGCGCCTGCCCTCCCCTGCCTGGGCACCGCCCGCGGG
:::
TTTGCCAGACCAAGAAGAAGCCATGTGCCCCAGCATCTACACTTCCTGTGCCTGGGCATCGTCCCCAGG
290      300      310      320      330      340      350

1320      1330      1340      1350      1360      1370      1380
GACGGCCCGCGACCGCAGCGGAGACAAGGACCTTCCTCGTTGGCCGCCCTCAGCGCTGGCCCTGGTGTG
:::
GACATCCCGAGAACGCAGTGGTGACAAGGACCTGCCCTCATTGGC-----TGTG
360      370      380      390      400

1390      1400      1410      1420      1430      1440      1450
GGGCTGTGTGAGGAGCATGGGTCTCCGGCAGCCCCCAGCACTTACTGGGCCAGGCCAGTTGCTGGCC
:::
GGCATATGTGAGGAGCATGGATCCGCCATGGCCCCCAGCACATCCTGGCCTCTGGCTCAACTGCTGGCC
410      420      430      440      450      460      470

1460      1470      1480      1490      1500      1510      1520
CTAAGTTGTACCCCAAACCTCTACACAGACATCCACACACACACA--CACACAC--TCTCACACACT
:::
CCAAGCTGTACCCCAAGCTATACACAGATGTGCACACACACACACATACACACACCTGCACTCACACGCT
480      490      500      510      520      530      540

1530      1540      1550      1560      1570      1580
CACACGT-GGAGGGCAAGGT-C-----CACCAGCATCCACTATCAGTGCTAGACGGCACCCTATCTGC
:::
CTCATGTTGGAGGGCAAGGTTTCATCAACACCAGCATGTCCACTATCAGTGCTAAA-TACAGCGAATCTCC
550      560      570      580      590      600      610

1590      1600      1610      1620      1630      1640      1650
AGTGGGCACGGGGGGGCCGGCCAGACAGGCAGACTGGGAGGATGGAGGACGGAGCTGCAGACGAAGGCAG
AA---GCACTGTGT-----CCTGA--GGTAGGCAT-----TTGGGGGCCAAGGCAACAG--GTTGG--G
620      630      640      650      660
```

FIG.28A

```

1660      1670      1680      1690      1700      1710      1720
GGGACCCATGGCGAGGAGGAATGGCCAGCACCCAGGCAGTCTGTGTGTGAGGCATAGCCCCTGGACACA
AGAATTGAGAAACAATGGAGGAAG--AGTATCTTAGGGTGCCT-TATGGTGGACA--CTCACAAACTTG
   670       680       690       700       710       720

1730      1740      1750      1760      1770      1780      1790
CACACACAGACACACACTGCCTGGA-TGCATGTATGCACACATGCGCGCACACGTGCTCCCTGAAG
GCCATATAGATGTATGTACTACAGATGAACAGCCAGCCAGATTACACACGCACATGTTTAAAC-GTGT
   730       740       750       760       770       780       790

1800      1810      1820      1830      1840      1850      1860
GCACACGTACGCA-CA-CACGCACATGCACAGATATGCCGCCTGGGCACACAGATAAGCTGCCCAAATGC
AAACGTGTGCACAACTGCACACACAA-C-CTGAGAAACCTTCAGGAGGATTGTGGTG-TGAC--TTTGC
   800       810       820       830       840       850       860

1870      1880      1890      1900      1910      1920      1930
ACGCACACGCA-CAGAGACATGCCAGAACATACAAGGACATG-CTGCCTGAACATA--CACACGCACACC
AGTGACATGTAGCGATGGCTAGTTGAAGGAATCTCCCTCATGTCTTAGTGGTCATGGCCACTTCCCACC
   870       880       890       900       910       920       930

1940      1950      1960      1970      1980      1990
CATGCGCAGATGTG--CTGCCTGGACACACACACACACGGATATGCTGTCTGGACGCACACACGTGC
CCTGCCCATCTGTGTTCTGCCTGGCCTTGGTGGTGCTTCCG--TGTGCC--CTGGGTTTTT-CAGGAAC
   940       950       960       970       980       990

2000      2010      2020      2030      2040      2050      2060
AGATATGGTATCCGGACACACACGTGCACAGATATGCTGCCTGGACACACAGATAATGCTGCCTTGACAC
C---CTATCAACCTGACTGGGGTGAGCA-----GTGCAGCCATGCNTGGAGGTTTGAGCCACC---CTC
   1000      1010      1020      1030      1040      1050

2070      2080
ACACATGCACGGATATTG
CC-CTTGCTAGAGAGAAG
   1060      1070

```

FIG.28B

```

      10      20      30      40      50      60      70
inputs MTPSPLLLLLLPLLLLGAFPPAAAARGPPKMAKVVPRQVARLGRTVRLQCPVEGDPPLTMWTKDGRTI
-----
      80      90     100     110     120     130     140
inputs HSGWSRFRVLPQGLKVQVEREDAGVYVCKATNGFGSLSVNYTLVVLDISPAGESLGPDSSSGGQEDPA
-----
     150     160     170     180     190     200     210
inputs SQQWARPRFTQPSKMRRRVVIARPVGSSVRLKCVASGHPRPDITWMKDDQALTRPEAAEPRKKKWTLSLKN
-----
     220     230     240     250     260     270     280
inputs LRPEDSGKYTCRVSNRAGAINATYKVDVIQRTSRKPVLTGTHPVNTTVDFGGTTSFQCKVRSVDVKPVIQW
      ...
      -----RVR-----
     290     300     310     320     330     340     350
inputs LKRVEYGAEGRHNSTIDVGGQKFVVLPTGDVWSRPDGSYLNKLLITRARQDDAGMYICLGANTMGYSFRS
      .....
      -----PTGDVWSRPDGSYLNKLLISRARQDDAGMYICLGANTMGYSFRS
              10      20      30      40
     360     370     380     390     400     410     420
inputs AFLTVLPDPKPPGPPVASSSSATSLPWPVVIGIPAGAVFILGTLWWLCQAQKKPCTPAPAPPLPGHRPP
      .....
      AFLTVLPDPKPPGPPMASSSSSTSLPWPVVIGIPAGAVFILGTVLLWLCQTKKKPCAPASTLPVPGHRPP
      50      60      70      80      90     100     110
     430     440     450     460     470     480
inputs GTARDRSGDKDLPSLAALSAGPGVGLCEEHGSAPAQHLLGPGPVAGPKLYPKLYTDIHTHTHTSHSTH-
      .....
      GTSRERSGDKDLPSLA------VGICEEHGSAMAPQHILASGSTAGPKLYPKLYTDVHTHTHTHTCTHT
      120     130     140     150     160     170     180
     490     500
inputs -----SHVEGKVHQHIHYQC
      ...
      LSCWRARFINTSMSTISAKYSESPSTVS
      190     200

```

FIG.29

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inputs GT-----
A
GTGTCACCGCCTCTGTGTCCCTCCTTCTCCTGGCTGTGGGCCTGCGGCTGGCTGGAACCTCAACCCCA
10 20 30 40 50 60 70

inputs -----
GTGATCCCAATACCTGCAGCTTCTGGGAAAGCTTCACTACCACCACCAAGGAGTCCCACTCCCGCCCTT
80 90 100 110 120 130 140

inputs -----
CAGCCTGCTCCCTCAGAGCCCTGCGAGCGGCCCTGGGAGGGCCCCATACTTGCCCCAGCCACAACT
150 160 170 180 190 200 210

inputs -----
CAGAGGAAACTCCTGGCTTCTAGGGATTCTTCTGCATGGTCTGTGTCGGGGCTGGAGTGCAGTGCCGAG
220 230 240 250 260 270 280

inputs -----
ATCGTAGTGCACTGCAACCTCAAACAGGGAATGCGCTTTCTATGCGCCCTCAGCCCAGAGTGTTGAGTGG
290 300 310 320 330 340 350

inputs -----
TGCCCCCTCCCTGGCCTCCCCTGGCCACACTGTGGTGGTGAAGACGGACCACCGCCAGCGCCTGCAGTGC
360 370 380 390 400 410 420

inputs -----
TGCCATGGCTTCTATGAGAGCAGGGGGTTCTGTGTCCCGCTCTGTGCCAGGAGTGTGTCCATGGCCGTT
430 440 450 460 470 480 490

inputs -----
GTGTGGCACCCAATCAGTGCCAATGTGTGCCAGGCTGGCGGGGCGACGACTGTTCCAGTGCCCCGAAGTGC
500 510 520 530 540 550 560

inputs -----
CCTTCAGCCCTGTACCCCTGGCTACTATGGCCCTGCCTGCCAGTTCGCTGCCAGTGCCATGGGGCACCC
570 580 590 600 610 620 630

inputs -----
TGCGATCCCAGACTGGAGCCTGCTTCTGCCCCGAGAGAGAACTGGGCCAGCTGTGACGTGTCCTGTT
640 650 660 670 680 690 700

FIG.30A

inputs
 CCCAGGGCAGCTTCTGGCTTCTTCTGCCCCAGCACCCATCCTTGCCAAAATGGAGGTGTCTTCCAAACCCC
 710 720 730 740 750 760 770

inputs
 ACAGGGCTCCTGCAGCTGCCCCCTGGCTGGATGGGCACCATCTGCTCCCTGCCCTGCCAGAGGGCTTT
 780 790 800 810 820 830 840

inputs
 CACGGACCCAAGTCTCCAGGAATGTGCTGCCACAACGGCGGCCTCTGTGACCGATTCACTGGGCAGT
 850 860 870 880 890 900 910

inputs
 GCCGCTGCGCTCCGGTTACACTGGGGATCGGTGCCGGGAGGAGTGCCCGGTGGGCCGCTTTGGGCAGGA
 920 930 940 950 960 970 980

inputs
 CTGTGCTGAGACGTGCGACTGCGCCCCGAGCGCCGTTGCTTCCGGCCAACGGCGCATGTCTGTGCGAA
 990 1000 1010 1020 1030 1040 1050

inputs
 CACGGCTTCACTGGGGACCGCTGCACGGATCGCTCTGCCCCGACGGCTTCTACGGTCTCAGCTGCCAGG
 1060 1070 1080 1090 1100 1110 1120

inputsCGACC.....
 CCCCCTGCACCTGCGACCGGGAGCACAGCCTCAGCTGCCACCGATGAACGGGGAGTGCTCCTGCCTGCC
 1130 1140 1150 1160 1170 1180 1190

inputs10
 CACGC.....
 GGGCTGGGCGGGCTCCACTGCAACGAGAGCTGCCCGCAGGACACGCATGGGCCAGGGTGCCAGGAGCAC
 1200 1210 1220 1230 1240 1250 1260

inputs
 TGTCTCTGCCTGCACGGTGGCGTCTGCCAGGCTACCAGCGGCTCTGTCACTGCGCGCCGGTTACACGG
 1270 1280 1290 1300 1310 1320 1330

inputs
 GCCCTCACTGTGCTAGTCTTTGTCTCCTGACACCTACGGTGTCAACTGTTCTGCACGCTGCTCATGTGA
 1340 1350 1360 1370 1380 1390 1400

FIG. 30B

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```

inputs .....
      AAATGCCATCGCCTGCTCACCCATCGACGGCGAGTGCGTCTGCAAGGAAGGTTGGCAGCGTGGTAACTGC
      1410      1420      1430      1440      1450      1460      1470

inputs .....
      TCTGTGCCCTGCCACCCGGAACCTGGGGCTTCAGTTGCAATGCCAGCTGCCAGTGTGCCCATGAGGCAG
      1480      1490      1500      1510      1520      1530      1540

inputs .....G
      TCTGCAGCCCCCAAACCTGGAGCCTGTACCTGCACCCCTGGGTGGCATGGGGCCCACTGCCAGCTGCCCTG
      1550      1560      1570      1580      1590      1600      1610

inputs TCCG.....GTGACCCCT
      TCCGAAGGGGAGTTTGGAGAAGGTTGTGCCAGTCGCTGTGACTGTGACCACTCTGATGGCTGTGACCCCT
      1620      1630      1640      1650      1660      1670      1680

inputs 30 40 50 60 70 80 90
      GTTCATGGACAGTGCCGATGTCAGGCTGGTTGGATGGGCACACGCTGCCACCTGCCTTGCCCGGAGGGCT
      1690      1700      1710      1720      1730      1740      1750
      GTTCATGGACGCTGTCACTGCCAGGCTGGCTGGATGGGTGCCCGCTGCCACCTGTCTCCCTGAGGGCT

inputs 100 110 120 130 140 150 160
      TTTGGGGAGCCAACTGCAGTAACACCTGTACCTGCAAGAATGGTGGTACCTGTGTGTCTGAGAATGGCAA
      1760      1770      1780      1790      1800      1810      1820
      TATGGGGAGTCAACTGTAGCAACACCTGCACCTGCAAGAATGGGGGACCTGTCTCCCTGAGAATGGCAA

inputs 170 180 190 200 210 220 230
      CTGCGTGTGCGCACCAGGTTCCGAGGGCCCTCCTGCCAGAGGCCCTGCCCGCCTGGTCGCTATGGCAAA
      1830      1840      1850      1860      1870      1880      1890
      CTGCGTGTGTGCACCCGATTCCGGGGCCCTCCTGCCAGAGATCCTGTGAGCCTGGCCGCTATGGCAAA

inputs 240 250 260 270 280 290 300
      CGCTGTGTGCAATGCAAGTGTAACAACAACCATTTCTTCTGCCACCCATCGGACGGGACCTGCTCCTGCC
      1900      1910      1920      1930      1940      1950
      CGCTGTGTGCCCTGCAAGTG---CGCTAACCACTCCTTCTGCCACCCCTCGAACGGGACCTGCTACTGCC

inputs 310 320 330 340 350 360 370
      TGGCGGGCTGGACAGGCCCTGACTGCTCCGAGGCATGTCCCCAGGCCACTGGGGACTCAAATGCTCCCA
      1960      1970      1980      1990      2000      2010      2020
      TGGCTGGCTGGACAGGCCCTGACTGCTCCAGCCATGCCCTCCAGGACACTGGGGAGAAAAGTGTGCCCA

inputs 380 390 400 410 420 430 440
      ACTCTGCCAGTGTATCATGGTGGGACCTGCCACCCCAAGGATGGGAGCTGTATCTGCACGCCAGGCTGG
      2030      2040      2050      2060      2070      2080      2090
      GACCTGCCAATGTACCATGGTGGGACCTGCCATCCCAAGGATGGGAGCTGTATCTGCCCCCTAGGCTGG

```

FIG.30C

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      450      460      470      480      490      500      510
inputs ACTCGACCCAACCTCCTTGAAGGCTGCCACCAAGAATGTTTGGTGCAACTGCTCCCAGCTATGTCAGT
      2100      2110      2120      2130      2140      2150      2160
      520      530      540      550      560      570      580
inputs GTGATCTCGGAGAGATGTGCCACCCAGAGACTGGGGCTTGTGTCTGTCCCCAGGACACAGTGGTGCAGA
      2170      2180      2190      2200      2210      2220      2230
      590      600      610      620      630      640      650
inputs CTGCAAAATGGGAAGCCAGGAGTCCCTTACCATAATGCCACCTCTCCCGTGACCCATAAATCACTGGGT
      2240      2250      2260      2270      2280      2290      2300
      660      670      680      690      700      710      720
inputs GCAGTGATTGGCATTGCAGTACTGGGAACCTCGTGGTGGCCCTGATAGCACTGTTTCATTGGCTACCGCC
      2310      2320      2330      2340      2350      2360      2370
      730      740      750      760      770      780      790
inputs AGTGGCAAAAGGGCAAGGAACATGAGCACTTGGCAGTGGCTTACAGCACTGGGCGGCTGGATGGCTCTGA
      2380      2390      2400      2410      2420      2430      2440
      800      810      820      830      840      850      860
inputs TTACGTCATGCCAGATGTCTCTCCGAGCTATAGTCACTACTCAACCCAGCTACCACACACTGTCT
      2450      2460      2470      2480      2490      2500      2510
      870      880      890      900      910      920      930
inputs CAGTGTCTCCTAACCCCCCGCCCCCTAACAAAGGTCCCAGGCAGCTCTTTGTGAGCTCTCAGGCCC
      2520      2530      2540      2550      2560      2570      2580
      940      950      960      970      980      990      1000
inputs CTGAGCGGCCAAGCAGAGCCACGGGCGTGAGAACCATAACCACTGCCCGCTGACTGGAAGCACCGCCG
      2590      2600      2610      2620      2630      2640      2650
      1010      1020      1030      1040      1050      1060      1160
inputs GGAGCCCCAT-----GACAGAGGCGCCAGCCACCTGGACCGAAGCTATAGCTGTAGCTATAGC
      2660      2670      2680      2690      2700      2710      2720
      1070      1080      1090      1100      1110      1120      1130
inputs CACAGGAATGGCCAGGACCATCTGTGCATAAAGGTCCCATCTCTGAAGAGGAGCTAGGGGCAAGCGTTA
      2730      2740      2750      2760      2770      2780
      -----AATGGCCAGGCCATTCTACGATAAAGGCTCATCTCTGAAGAGGAGCTCGGGCCAGTGTGG

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FIG.30D

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      1140      1150      1160      1170      1180      1190      1200
inputs TGTCCCTGAGCAGTGAGAACCCCTATGCTACCATCCGAGACCTGCCAGCCTGCCTGGGGAACCCCGAGA
      .....
      CTTCCCTGAGCAGTGAGAACCCATATGCCACCATCCGGGACCTGCCAGCTTGCCAGGGGGCCCCGGGA
2790      2800      2810      2820      2830      2840      2850

      1210      1220      1230      1240      1250      1260      1270
inputs AAGTGGCTATGTGGAGATGAAAGGACCTCCATCAGTGTCCCCTCCCAGGCAGTCTCTTCATCTCCGGGAC
      ...
      GAGCAGCTACATGGAGATGAAAGGCCCTCCCTCAGGATCTGCCCCAGGCAGCCTCCTCAGTTTTGGGAC
2860      2870      2880      2890      2900      2910      2920

      1280      1290      1300      1310      1320      1330      1340
inputs AGGCAG---CAGCGGCAACTGCAGCCACAGAGGGACAGCGGCACCTATGAGCAGCCAGCCCCCTTGAGCC
      ::::
      AGCCAGAGGCGGCGGCAACCCAGCCACAGAGAGACAGTGGCACCTACGAGCAGCCAGCCCCCTGATCC
2930      2940      2950      2960      2970      2980      2990

      1350      1360      1370      1380      1390      1400      1410
inputs ATAATGAAGAGTCTTTGGGCTCCACGCCCCCGCTTCTCCAGGCCTGCCTCCTGGTCACTACGACTCCCC
      ....
      ATGACCGAGACTCTGTGGGCTCCAGCCCCCTCTGCCTCCGGGCCTACCCCCGGCCACTATGACTCACC
3000      3010      3020      3030      3040      3050      3060

      1420      1430      1440      1450      1460      1470      1480
inputs CAAGAACAGCCATATCCCTGGACACTATGACTTGCTCCAGTACGGGATCCTCCATCCCCTCCATCCCGG
      .....
      CAAGAACAGCCACATCCCTGGACATTATGACTTGCTCCAGTACGGGATCCCCATCAGCTCCACTTCGA
3070      3080      3090      3100      3110      3120      3130

      1490
inputs CGCCAGGACCGC
      .....
      CGCCAGGACCGT
3140      3150
```

FIG.30E

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1890 1900 1910 1920 1930 1940 1950
GACCACTCTGATGGCTGTGACCCCTGTTTCATGGACGCTGTCAGTGCCAGGCTGGCTGGATGGGTGCCCCGCT
GACC-CAC-GCGTCCGGTGACCCCTGTTTCATGGACAGTGCCGATGTCAGGCTGGTTGGATGGGCACACGCT
10 20 30 40 50 60 70
1960 1970 1980 1990 2000 2010 2020
GCCACCTGTCTGCCCTGAGGGCTTATGGGAGTCAACTGTAGCAACACCTGCACCTGCAAGAATGGGGG
GCCACCTGCCTTGCCCGGAGGGCTTTTGGGAGCCAACTGCAGTAACACCTGTACCTGCAAGAATGGTGG
80 90 100 110 120 130 140
2030 2040 2050 2060 2070 2080 2090
CACCTGTCTCCCTGAGAATGGCAACTGCGTGTGTGCACCCGGATTCCGGGGCCCTCCTGCCAGAGATCC
TACCTGTGTGTCTGAGAATGGCAACTGCGTGTGCGCACCCAGGGTTCCGAGGCCCTCCTGCCAGAGGCC
150 160 170 180 190 200 210
2100 2110 2120 2130 2140 2150 2160
TGTACGCTGGCCGCTATGGCAAACGCTGTGTGCCCTGCAAGTG--CGCTAACCACTCCTTCTGCCACC
TGCCCGCCTGGTCTGCTATGGCAAACGCTGTGTGCAATGCAAGTGTAAACAACAACCACTTCTTCTGCCACC
220 230 240 250 260 270 280
2170 2180 2190 2200 2210 2220 2230
CCTCGAACGGGACCTGCTACTGCTGCTGGCTGGACAGGCCCGACTGCTCCAGCCATGCCCTCCAGG
CATCGGACGGGACCTGCTCCTGCTGGCGGGCTGGACAGGCCCTGACTGCTCCGAGGCATGTCCCCCAGG
290 300 310 320 330 340 350
2240 2250 2260 2270 2280 2290 2300
ACACTGGGGAGAAACTGTGCCAGACCTGCCAATGTACCATGGTGGGACCTGCCATCCCCAGGATGGG
CCACTGGGGACTCAAATGCTCCCAACTCTGCCAGTGTTCATCATGGTGGGACCTGCCACCCCAAGGATGGG
360 370 380 390 400 410 420
2310 2320 2330 2340 2350 2360 2370
AGCTGTATCTGCCCCCTAGGCTGGACTGGACACCACTGCTTAGAAGGCTGCCCTCTGGGGACATTTGGTG
AGCTGTATCTGCACGCCAGGCTGGACTGGACCCAACCTGCTTGAAGGCTGCCACCAAGAATGTTTGGTG
430 440 450 460 470 480 490
2380 2390 2400 2410 2420 2430 2440
CTAACTGCTCCCAGCCATGCCAGTGTGGTCTGGAGAAAAGTGCCACCCAGAGACTGGGGCTGTGTATG
TCAACTGCTCCCAGCTATGTCAAGTGTGATCTCGGAGAGATGTGCCACCCAGAGACTGGGGCTGTGTCTG
500 510 520 530 540 550 560
2450 2460 2470 2480 2490 2500 2510
TCCCCAGGGCACAGTGGTGCACCTTGCAGGATTGGAATCCAGGAGCCCTTTACTGTGATGCCGACCACT
TCCCCAGGACACAGTGGTGCAGACTGCAAAATGGGAAGCCAGGAGTCTTACCAATAATGCCACCTCT
570 580 590 600 610 620 630
2520 2530 2540 2550 2560 2570 2580
CCAGTAGCGTATAACTCGTGGGTGCAGTGATTGGCATTGCAGTGCTGGGGTCCCTTGTGGTAGCCCTGG
CCCGTGACCCATAACTCACTGGGTGCAGTGATTGGCATTGCAGTACTGGGAACCCCTCGTGGTGGCCCTGA
640 650 660 670 680 690 700

FIG.31A

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```

      2590      2600      2610      2620      2630      2640      2650
TGGCACTGTTTCATTGGCTATCGGCACTGGCAAAAAGGCAAGGAGCACCACCACCTGGCTGTGGCTTACAG
TAGCACTGTTTCATTGGCTACCGCCAGTGGCAAAAAGGCAAGGAACATGAGCACTTGGCAGTGGCTTACAG
      710      720      730      740      750      760      770

      2660      2670      2680      2690      2700      2710      2720
CAGCGGGCGCCTGGACGGCTCCGAGTATGTCATGCCAGATGTCCCTCCGAGCTACAGTCACTACTACTCC
CACTGGGCGGCTGGATGGCTCTGATTACGTCAAGCCAGATGTCTCTCCGAGCTATAGTCACTACTACTCC
      780      790      800      810      820      830      840

      2730      2740      2750      2760      2770      2780
AACCCAGCTACCACACCCTGTGCGAGTGTCCCCAAACCCCAACCCCTAACAGGTTCCAGGC---C
AACCCAGCTACCACACACTGTCTCAGTGTCTCTCTAAACCCCGCCCTAACAGGTTCCAGGCACTC
      850      860      870      880      890      900      910

2790      2800      2810      2820      2830      2840      2850
CGCTCTTTGCCAGCCTGCAGAACCTGAGCGGCCAGGTGGGGCCCAAGGGCATGATAACCACACCACCCT
AGCTCTTTGTAGCTCTCAGGCCCTGAGCGGCCAAGCAGAGCCACGGGCGTGAGAACCATACCACT
      920      930      940      950      960      970      980

2860      2870      2880      2890      2900      2910      2920
GCCTGCTGACTGGAAGCACCGCGGGAGCCCTCCAGGGCCTCTGGACAGGGGAGCAGCGCCTGGAC
GCCGCTGACTGGAAGCACCGCGGGAGCCCAT-----GACAGAGGCGCCAGCCACCTGGAC
      990      1000      1010      1020      1030

2930      2940      2950      2960      2970      2980      2990
CGAAGCTACAGCTATAGCTACAGC-----AATGGCCCAGGCCATTCTACGATAAAGGGCTCATCTCTG
CGAAGCTATAGCTGTAGCTATAGCCACAGGAATGGCCCAGGACCATCTGTCTATAAGGTCCCATCTCTG
      1040      1050      1060      1070      1080      1090      1100

      3000      3010      3020      3030      3040      3050      3060
AAGAGGAGCTCGGGGCCAGTGTGGCTTCCCTGAGCAGTGAGAACCCATATGCCACCATCCGGGACCTGCC
AAGAGGGAAGTGGGGCAAGCGTTATGTCCCTGAGCAGTGAGAACCCCTATGCTACCATCCGAGACCTGCC
      1110      1120      1130      1140      1150      1160      1170

      3070      3080      3090      3100      3110      3120      3130
CAGCTTGCCAGGGGGCCCCGGGAGAGCAGCTACATGGAGATGAAAGGCCCTCCCTCAGGATCTGCCCCC
CAGCCTGCCTGGGGAACCCCGAGAAAGTGGCTATGTGGAGATGAAAGGACCTCCATCAGTGTCCCCTCCC
      1180      1190      1200      1210      1220      1230      1240

      3140      3150      3160      3170      3180      3190      3200
AGGCAGCCTCCTCAGTTTTGGGACAGCCAGAGCGGGCGGCAACCCAGCCACAGAGAGACAGTGGCACCT
AGGCAGTCTCTTCATCTCCGGGACAGGCAG---CAGCGGCAACTGCAGCCACAGAGGGACAGCGGCACCT
      1250      1260      1270      1280      1290      1300      1310

      3210      3220      3230      3240      3250      3260      3270
ACGAGCAGCCAGCCCCCTGATCCATGACCGAGACTCTGTGGGCTCCAGCCCCCTCTGCCTCCGGGCT
ATGAGCAGCCAGCCCCCTGAGCCATAATGAAGAGTCTTTGGGCTCCAGCCCCCGCTTCTCCAGGCCT
      1320      1330      1340      1350      1360      1370      1380

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FIG.31B

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      3280      3290      3300      3310      3320      3330      3340
ACCCCCGGGCACTATGACTCACCCAAGAACAGCCACATCCCTGGACATTATGACTTGCCTCCAGTACGG
      1390      1400      1410      1420      1430      1440      1450
GCCTCCTGGTCACTACGACTCCCCAAGAACAGCCATATCCCTGGACACTATGACTTGCCTCCAGTACGG

      3350      3360      3370      3380      3390      3400      3410
CATCCCCATCACCTCCACTTCGACGCCAGGACCGTTGAGGAGCCAGGATGGTATGGCAGAGGCCAGCAC
      1460      1470      1480      1490      1500      1510
CATCCTCCATCCCCTCCATCCCGGCCAGGACCGCTGAAGAGCCGGCATGGTATG---GGAGC-----

      3420      3430      3440      3450      3460      3470      3480
ACCTGGCTGTTGCTGCTCAAGGCTGGGGACAGAGCCTAGTGTACCCCTGCCAGGAGCAGGGAGTGGACCG
      1520      1530      1540      1550
-----GTGCCCTA-TGTACCT-TGCCAGGAGCAGGGACTGGACCA

      3490      3500      3510      3520      3530      3540      3550
GCAGGCTGTGAACATGAACAACGCTTAACAGAGCAAGTGATGG-GAGCCTTGTTCTGGG-TTCTACCAT
      1560      1570      1580      1590      1600      1610      1620
GCAGGCCACGAACAGAAACA---CTTGGTGAAGTGAACAGAGACGGACTGTGGCCCTGTGCTTCCACCGA

      3560      3570      3580      3590      3600      3610
GGGAGACGCTGATCAGCAGGATGCCTGGCTCCCTTTCCCAACCCACTGCTCCCAAGGCCTCCAGGGC---
      1630      1640      1650      1660      1670      1680
GGGAGACACTAGTTGACAAAGTGTCTAACCCCTCTTTTCCAAACCCACTGCT--CAAGTCCCTGTGGACATA

      3620      3630      3640      3650      3660      3670      3680
--CCTGTGTACATAAACTGGTGGGTTGGAAGTTGCTGGGTAACTCTGATTTCAGACATGCGTGTGGGGT
      1690      1700      1710      1720      1730      1740      1750
AGCTGGTGGGCAGAATGTTGTTGTACAAGTGTGATTTTAGATCGATTTTTTTAAAGTATGTGTTGGGT

      3690      3700      3710      3720      3730      3740      3750
ACCTTTTCTGTGC--ATGCTCAGCCTGGGCTCTGTGCGTGTGTGTTTCTGTGATTTTAGAAGGGTACC
      1760      1770      1780      1790      1800      1810      1820
ACCTTTTCTGTGTGTATGCTCAGGCAGG--CTGTG---TGTGTCTCTAGTTGGCTTTAGAGGGAGTCA

      3760      3770      3780      3790      3800      3810      3820
AG-GCAGGTTCTGTCTAGGGCACTTACCATTAGTAGGGAGATGGAACCAACCAATTAACCTCTAGCAA
      1830      1840      1850      1860      1870      1880
GGTATAGGTTCTG-CTTCTGCACTTTCCATCTTATCTAGTAGTCAG--CTTCCAAGCTTA-CTAGTTA

      3830      3840      3850      3860      3870      3880      3890
TAGCCTCCTAACTGGCCTCCTCCATTGATTGAGTGAACCTTCCAATGCATGGCTCATAATTTCAAAATAC
      1890      1900      1910      1920      1930
GAGC-TCCA-----CCAGCAGCA--GGCCCTAACTACCTGCCT-----GCCC-----TTCA-----C

      3900      3910      3920      3930      3940      3950      3960
AGGCTGGTTAGTTACTCCCTACCTGAAAGCCTTCATAGGTGCCTCTTTGCTCTTCTGCCAGTATCAAAAC
      1940      1950      1960      1970
---CCAGTAA--TCCTCCATGTCT--TTGC--TCAGAGGA-----TTGCTC-----CC-----CGACTC

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FIG 31C

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3970      3980      3990      4000      4010      4020      4030
TTTTGAAGGCCTTAAAGGCCCTGCTTTGCCTGGCCCATCTGTCTCTCCAGCCTCACCTTGAACCTGTGTTT
1980      1990      2000      2010      2020
TGGTGTGTGCTCT-----CCTGGTACGCCTTG----ACGGTC-CTGCAGTCTC-CCT-----TTC

4040      4050      4060      4070      4080      4090      4100
CTGTCACTGCACGCCAGTCACACCGGCCTCTAGGTCTCTGTAGGCCACTCTTCTTTCTGGCACAGGGA
2030      2040      2050      2060      2070      2080
CCGTCT-TGCT--TCATTCTTTC--CCAGAATGAAGGC-TGTCTGCCACCCTACTTCCCAGCCCAGGAA

4110      4120      4130      4140      4150      4160      4170
CCTGCACACCTGGAGTGCCCTTCTCCCCACTCGCTGTTACCCCTGCTTTTCTTTACACCTCCTCC
2090      2100      2110      2120      2130      2140      2150
TTGGCACATCTAAGTT--CAGCCTTCTTAAGTTACCCGTTGAGTCTGCTTGCCTT--CACATATTCC

4180      4190      4200      4210      4220      4230      4240
TCAGGGAAGTGCCACCCCTCCGTACATCTTTCACAGCCCTGATTGCAGCTGTGTTCACTCACCAGGTACC
2160      2170      2180      2190      2200
ACAGAAC--CCCACC--CC--ACATCT--GCTTC--ATAGCTACTCTCTTCTC-CAC--GTACC

4250      4260      4270      4280      4290      4300      4310
TGCAGAAGGCCTACAGGTGCCAGGCACCTTTTAAATGGGTTCTTTCTTTATGTGATTATTTGATTAAATC
2210      2220      2230      2240      2250      2260      2270
CACAGAAGGCAGAAAGTGGTACCAGGCAAGA--AGATGGGATTGTTGCATTTTGT--TTTGTTTTGGAGAC

4320      4330      4340      4350      4360      4370      4380
TCTGCCTCCCCACTAGACTGTAAGCTCCCTGAAGGCAAGAATCCTG--TGCTTATGCTCAATATTAGCT
2280      2290      2300      2310      2320      2330      2340
TCTGTCTCACTATGTAGTCTGCTGGCTGGCACTCAAGAGCTCTGCCTGCCTCTGCTCTTGAAGTCT

4390      4400      4410      4420      4430      4440
CTCCCTT--GGCACAGAGT--AGGCACTCAACAAA-TGCTCCCCAAAAGGCTGAGTGGCTGACTGAATT
2350      2360      2370      2380      2390      2400
GGGTTTAAACGGCTCAGGGTCACATGCACAGCTCAAGCTGCACTCCGATGTGCT--TTCC--CCTGTTGC

4450      4460      4470      4480      4490      4500      4510
AAGTACCAGTGACATGCAGTAACTGCTAAGATAGATGAGCCATCTGTATGCTCTGACAGTTACAG-ACTG
2410      2420      2430      2440      2450      2460
TAGAT---TAGCGT-CTGCCTCCCCCTAG-TGGAGAGGCTGATCGCCAGCTCT--CTGATGCAGGACTC

4520      4530      4540      4550      4560      4570      4580
AATAAGTTGGAGACT-TCCCTAAAGGGTGGCATTTCCTCAGGGTAACAACGCAGAGCTCAGGTGTGGGAA
2470      2480      2490      2500      2510      2520      2530
TGGTGTTTAGGCTCACTCACTATTGTTT-CCTTGGCACAGGGTAGTCACTCAATA--AATGTTCTCTTA

4590      4600      4610
GGTGCCAGGGGCAGGGGTGCAGAGGGGCTGAGGC
2540      2550      2560
AAAGCTGAAAAAAAAAAAAAAAAAGGGCGGCCGC

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FIG. 31D

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      10      20      30      40      50      60      70
inputs  MSPPLCPLLLLAVGLRLAGTLNPSDPNTCSFWE$FTTTTKESH$RPF$LLPSEP$CERPWE$GPH$TC$P$P$QT
      .....

      80      90      100     110     120     130     140
inputs  QRKLLASRDSFCMVCV$GAGVQWRDR$SALQPQTGNAL$MRPQPRVLSGAPSLASP$GHTVVVKTDHRQRLQC
      .....

      150     160     170     180     190     200     210
inputs  CHGFY$SRGFCVPLCAQ$ECVHGRCVAPNQ$CQCVPGWRGDDC$SAPNCLQPCTPGYYG$PACQFRCQCHGAP
      .....

      220     230     240     250     260     270     280
inputs  CDPQTGACFCPAERTGP$SCDVSCSQGTS$GFFC$P$THPCQNGGVFQTPQG$SC$CPPGWMGTIC$SLPCPEGF
      .....

      290     300     310     320     330     340     350
inputs  HGPNC$QE$CRCHNGGLCDRFTGQCR$CAPGYTGDR$CREECPVGRFGQCAETCD$CAPDARCFPANGACLCE
      .....

      360     370     380     390     400     410     420
inputs  HGFTGDRCTDRLCPDGFYGL$SCQAPCTCDREH$LSCHPMNGECSCLPGWAGLHCNE$SCPQDTHGPGCQE$H
      .....
                                     $THASG$
      .....

      430     440     450     460     470     480     490
inputs  CLCLHGGVCQAT$SGLCQCAPGYTGPH$CASLCP$PDYGVNCSARCSCENAIAC$SPIDGECVCKEGWQR$GNC
      .....

      500     510     520     530     540     550     560
inputs  SVPCPPGTWGF$SCNASCQCAHEAVC$SPQTGACTCTPGWHGAHCQLPCPKGQFGE$G$CASRCD$CDHSDG$CDP
      .....
                                     $DP
      .....

      570     580     590     600     610     620     630
inputs  VHGR$CQ$CQAGW$MGARCHL$SCPEGLWGVNCSNTCTCKNGGTCLPEN$GNCVCAPGFRG$P$SCQ$R$SCQ$P$GRY$GK
      .....
VHGQCR$CQAGW$MGTRCHLPCPE$GFWGANC$NTCTCKNGGT$CVSENGNCVCAPGFRG$P$SCQ$R$PCPP$GRY$GK
      .....
      10      20      30      40      50      60      70

```

FIG. 32A

```

        640      650      660      670      680      690
inputs RCVPCCKAN-HSFCHPSNGTCYCLAGWTGPDSCQPCPPGHWGENCAQTCQCHHGGTCHPQDGSICPLGW
      ..... : : ..... : ..... : ..... : ..... : :
      RCVQCKCNNHSSCHPSDGTCSCLAGWTGPDSEACPPGHWGLKCSQLCQCHHGGTCHPQDGSICTPGW
      80      90      100     110     120     130     140

        700      710      720      730      740      750      760
inputs TGHHCLEGCPGLTGFAGNCSPQCQCGPGEKCHPETGACVCPGHSAPCRIGIEPFTVMPPTTPVAYNSLG
      ..... : : ..... : ..... : ..... : ..... : :
      TGPNCLEGCPPRMFGVNCSQLCQCDLGEMCHPETGACVCPGHSADCKMGSESFTIMPTSPVTHNSLG
      150     160     170     180     190     200     210

        770      780      790      800      810      820      830
inputs AVIGIAVLGSLVVALVALFIGYRHWQKGKEHHHLAVAYSSGRLDGSEYVMPDVPPSYSHYYSNPSYHTLS
      ..... : : ..... : ..... : ..... : ..... : :
      AVIGIAVLGTLVVALIALFIGYRQWQKGKEHEHLAVAYSTGRLDGSDYVMPDVSPSYSHYYSNPSYHTLS
      220     230     240     250     260     270     280

        840      850      860      870      880      890      900
inputs QCSPNPPPPNKVPGP-LFASLQNPERRPGGAQGHDNHTTLPADWKHRREPPPGPLDRGSSRLDRSYSYSYS
      ..... : : ..... : ..... : ..... : ..... : :
      QCSPNPPPPNKVPGSQLFVSSQAPERPSRAHGRENHTTLPADWKHRREPH---DRGASHLDRSYSCSYS
      290     300     310     320     330     340     350

        910      920      930      940      950      960      970
inputs --NGPGPFYDKGLISEEELGASVASLSSENPHYATIRDLPSLPGGPRESSYMEMKGPPSGSAPRQPPQFWD
      ..... : : ..... : ..... : ..... : ..... : :
      HRNGPGPFCHKGPISEEGLGASVMSLSSENPHYATIRDLPSLPGEPRESGYVEMKGPPSVSPRQSLHLRD
      360     370     380     390     400     410     420

        980      990      1000     1010     1020     1030     1040
inputs SQRRRQPQPQRDSGTYEQPSPLIHDRDSVGSQPPLPPGLPPGHYDSPKNSHIPGHYDLPPVRHPPSPPLR
      ..... : : ..... : ..... : ..... : ..... : :
      RQQR-QLQPQRDSGTYEQPSPLSHNEESLGSTPPLPPGLPPGHYDSPKNSHIPGHYDLPPVRHPPSPPSR
      430     440     450     460     470     480     490

        1050
inputs RQDR
      : : :
      RQDR

```

FIG.32B

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Input file T272Atrxa6b6; Output File T272Atrxa6b6.pat
Sequence length 3567

GTCCGACCCACGCGTCCGAGCCACACCCTGAAGGTGGTTGGAAGGAGGAAGGATCTAGGTCTGAGCACTGGAATTCC 79
CCAGAACAGCATCTGGCTTCCCAGACCCATGCTGGCCACCACTGATGTGTCCTTCCGGCTGCTGGCTGCAGTGTCTTC 158
TGTTGTTGGGTGCCCTGTGGCAGGCTGTGCAATGCCACTCTGTCCCCTCCTCCTCTGGCCCTAGGCCCTGCGTCTGGC 237
TGGAACACTCAACTCCAATGATCCCAATGTCTGTACCTTCTGGGAAAGCTTCACCAGACCACTAAGGAGTCCCACCTT 316
CGCCCCCTTCAGCCTGCCCCCAGCCGAGTCTGCGACAGGCCCTGGGAAGACCCCCACACCTGCGCTCAGCCTACGGTTG 395
TCTACCGGACTGTGTACCGTCAGGTGGTGAAGATGGACTCCCGCCACGCGCTGCAGTGTGTGGGGTTACTACGAGAG 474
CAGTGGAGCCTGTGTCCCACTCTGTGCCCAGGAGTGTGTCCACGGTCGCTGTGTGGCTCCTAATCGGTGCCAGTGTGCA 553
CCAGGCTGGCGGGGTGACCACTGTTCCAGTGAGTGTGCTCCTGGAATGTGGGGACCACAGTGTGACAGGCTCTGCCTCT 632
GTGGCAACAGCAGTTCTGTGATCCCAGGAGTGGGGTGTGTTTTTGCCTCTGGCTGCAGCCCCCGACTGCCTTCA 711
GCCTTGCCCCGATGGCCACTATGGTCCTGCCCTGCCAGTTTGATTGCCATGCTATGGGGCATCCTGTGACCCCCGGGAT 790
GGAGCCTGCTTCTGCCCCCAGGGAGAACAGGACCCAGGGCACTGATGGCTTCTTCTGCCCCAGAACTTATCCTTGCCA 869

M G V I C S 6
AAATGGAGGTGTTCTCAGGGCTCTCAAGGCTCCTGCAGTGTCCACCGGGCTGG ATG GGT GTC ATC TGT TCC 942

L P C P E G F H G P N C T Q E C R C H N 26
CTG CCA TGC CCA GAG GGT TTC CAC GGA CCC AAC TGT ACT CAG GAA TGT CGT TGC CAC AAT 1002

G G L C D R F T G Q C H C A P G Y I G D 46
GGT GCC CTT TGT GAC AGG TTT ACT GGG CAG TGC CAC TGT GCT CCT GGC TAT ATC GGG GAT 1062

R C R E E C P V G R F G Q D C A E T C D 66
CGG TGC CGT GAA GAG TGC CCT GTG GGC CGC TTC GGT CAA GAC TGT GCT GAG ACC TGT GAC 1122

C A P G A R C F P A N G A C L C E H G F 86
TGT GCT CCT GGC GCT CGT TGC TTT CCT GCC AAT GGC GCG TGT CTG TGC GAA CAT GGC TTC 1182

FIG.33A

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T G D R C T E R L C P D G R Y G L S C Q	106
ACA GGC GAC CGC TGC ACT GAG CGA CTC TGT CCA GAT GGC CGC TAT GGT CTG AGC TGC CAA	1242
D P C T C D P E H S L S C H P M H G E C	126
GAT CCC TGC ACC TGC GAC CCA GAA CAC AGT CTC AGC TGC CAC CCA ATG CAC GGC GAG TGC	1302
S C Q P G W A G L H C N E S C P Q D T H	146
TCC TGC CAG CCA GGT TGG GCG GGC CTC CAC TGC AAC GAG AGC TGC CCT CAG GAC ACG CAC	1362
G A G C Q E H C L C L H G G V C L A D S	166
GGA GCC GGT TGC CAG GAG CAC TGC CTC TGT CTG CAC GGC GGT GTT TGC CTC GCC GAC AGC	1422
G L C R C A P G Y T G P H C A N L C P P	186
GGC CTC TGC CCG TGT GCA CCT GGC TAC ACG GGA CCT CAC TGC GCT AAT CTT TGT CCA CCT	1482
N T Y G I N C S S H C S C E N A I A C S	206
AAC ACT TAT GGG ATC AAC TGT TCC TCC CAC TGC TCC TGT GAA AAT GCC ATT GCC TGC TCT	1542
P V D G T C I C K E G W Q R G N C S V P	226
CCT GTC GAC GGC ACG TGC ATC TGC AAG GAA GGT TGG CAG CGT GGT AAC TGC TCT GTG CCC	1602
C P P G T W G F S C N A S C Q C A H E G	246
TGT CCC CCT GGC ACC TGG GGC TTC AGT TGC AAT GCC AGT TGC CAG TGT GCC CAC GAG GGA	1662
V C S P Q T G A C T C T P G W R G V H C	266
GTC TGC AGC CCC CAA ACT GGA GCC TGT ACT TGC ACC CCT GGG TGG CGT GGG GTT CAC TGC	1722
Q L P C P K G Q F G E G C A S V C D C D	286
CAA CTT CCG TGC CCG AAG GGA CAG TTT GGT GAA GGT TGT GCC AGT GTC TGT GAC TGT GAC	1782
H S D G C D P V H G H C R C Q A G W M G	306
CAC TCC GAT GGC TGT GAC CCT GTT CAT GGA CAC TGC CGA TGT CAG GCT GGC TGG ATG GGC	1842
T R C H L P C P E G F W G A N C S N A C	326
ACA CGT TGC CAC CTG CCT TGC CCA GAG GGC TTT TGG GGA GCC AAC TGC AGC AAT GCC TGT	1902
T C K N G G T C V P E N G N C V C A P G	346
ACC TGC AAG AAT GGT GGC ACT TGT GTA CCT GAG AAC GGC AAC TGT GTG TGC GCA CCA GGC	1962

FIG.33B

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F R G P S C Q R P C P P G R Y G K R C V	366
TTC AGA GGC CCC TCC TGC CAG AGG CCC TGC CCG CCT GGT CGC TAT GGC AAA CGC TGT GTG	2022
P C K C N N H S S C H P S D G T C S C L	386
CCC TGC AAG TGC AAC AAC CAT TCT TCC TGC CAC CCG TCG GAT GGG ACC TGC TCC TGC CTG	2082
A G W T G P D C S E S C P P G H W G L K	406
GCA GGC TGG ACA GGC CCT GAC TGC TCT GAA TCA TGT CCC CCA GGC CAC TGG GGA CTC AAA	2142
C S Q P C Q C H H G A T C H P Q D G S C	426
TGC TCC CAA CCC TGC CAG TGT CAT CAT GGT GCC ACC TGC CAC CCC CAG GAT GGG AGC TGT	2202
V C I P G W T G P N C S E G C P S R M F	446
GTC TGC ATC CCA GGC TGG ACT GGA CCC AAC TGC TCG GAA GGC TGC CCA TCA AGA ATG TTT	2262
G V N C S Q L C Q C D P G E M C H P E T	466
GGT GTC AAC TGC TCC CAG CTA TGT CAG TGT GAT CCT GGA GAG ATG TGC CAC CCA GAG ACT	2322
G A C V C P P G H S G A H C K V G S Q E	486
GGG GCT TGC GTC TGT CCC CCA GGA CAC AGT GGT GCG CAC TGC AAA GTG GGC AGC CAG GAG	2382
S F T I M P T S P V I H N S L G A V I G	506
TCC TTC ACC ATA ATG CCC ACC TCT CCT GTG ATC CAT AAC TCA CTG GGT GCC GTG ATT GGC	2442
I A V L G T L V V A L V A L F I G Y R H	526
ATT GCA GTG CTG GGG ACC CTT GTG GTG GCC CTG GTA GCA CTG TTT ATT GGC TAC CGA CAC	2502
W Q K G K E H E H L A V A Y S T G R L D	546
TGG CAA AAG GGC AAG GAA CAT GAG CAC TTG GCA GTG GCT TAC AGC ACT GGG CGA CTG GAT	2562
G S D Y V M P D V S P S Y S H Y Y S N P	566
GGC TCC GAT TAC GTC ATG CCA GAT GTC TCT CCG AGC TAC AGT CAC TAC TAT TCC AAC CCT	2622
S Y H T L S Q C S P N P P P P N K I P G	586
AGC TAC CAC ACA CTG TCT CAG TGT TCT CCT AAC CCT CCA CCC CCT AAC AAG ATT CCA GGC	2682
S Q L F V S S Q A S E R P N R N H G R D	606
AGT CAG CTG TTT GTC AGC TCC CAG GCA TCT GAG CGG CCA AAC AGA AAC CAT GGG CGA GAT	2742

FIG.33C

N H A T L P A D W K H R R E S H D R A F	626
AAC CAC GCC ACA CTG CCC GCT GAC TGG AAG CAC CGA CCG GAG TCC CAT GAC AGA GCT TTC	2802
L R H Q P P G P K V *	637
CTC AGG CAC CAG CCA CCT GGA CCG AAG GTA TAG	2835
CTGTAGCTATGGCCACAGGAATGGCCCGGGCCATTCTGTCATAAAGGTCCCATCTCTGAAGAAGGACTAGGGGCAAGC	2914
GTTATGTCCCTGAGCAGTGAGAACCCTATGCGACCATCCGAGACCTGCCCGGCTGCCTGGGGAACCCCGAGAAAGCA	2993
GCTATGTGGAGATGAAAGGCCCTCCATCAGTGTCTCCCCCAGGCAGCCTCTTCATCTCCGGACAGGCAGCAGCAGCA	3072
ACTGCAGTCTCAGAGAGACAGCGGCACCTATGAGCAGCCCACTCCCTTGAGCCGTAATGAAGAGTCTGTGGGCTCCATG	3151
CCCCCTCTTCTCCGGGCTGCCACCCGGCCACTATGACTCGCCCAAAACAGCCACATCCCTGGACACTATGACTTGC	3230
CTCCAGTACGGCATCCTCCATCACCTCCATCCCGGCCAGGACCGCTGAGGAGCCAGCATGGTATGGGAGAGTGCCTG	3309
TGAACCTGCCAGGAGCAGGGCCTGGACCAGCAGGCCATGAATAGACATACTTGGTGAAGTGAACGAGACTGAGGATG	3388
GCTCTGCTTCCACCGAGGAGACACTAGTTGGCAAAGTGTCTAACCTCCCTTTTCCAGCCCATTGCTCAAGTCCCCCAG	3467
GCTGTGGACATGAGCTGGTGGGCAGAATGTTGTTGTTGAAGTCTGATTTTAGATTGATTTTTTAAAAAAAAAAAAAAAA	3546
AAAAAAAAAAGGCGGCCGC	3567

FIG.33D

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	10	20	30	40	50	60
human	GTC-GACCCACGCGTCCGCTCGAAGCGGGGACCCTCGCCCCGTCTCGGCTGTCCAGTCTCTCTCTCGC					
rat	GTCCGACCCACGCGTCCG-----AGC-----CACACCCTGAAGGTGTTGGAAGG-----					
	10	20	30	40		
	70	80	90	100	110	120
human	AGACCCCGGCGGTTCTACCCAGGCCGAGGGGAGACGGTGCCCCAAGGCAGGCTTCATA--TCCTGAA					
rat	AGG---GAAGGATCTAGGTCCTGAGCACTGG-----AATTCGCCAGAACAG-CATCTGGCTTCCAGA					
	50	60	70	80	90	100
	140	150	160	170	180	190
human	CGCTGG-GATCCCCCA-GGACATTCCCTGGCCCCAGGCCCCAGGTCCCAGGCCCCAGGGCTGAGCTGTG					
rat	CCCATGCTGGCCACCACTGATGTGTCCTT---CCGG---CTG---CTGGCTGCAGTCTGTTCTGTT					
	110	120	130	140	150	160
	210	220	230	240	250	260
human	GGCAGGCCCCACCTGGCCTCTGCAATGTCACCGCCTCTGTGTCCCTCTTCTCTGGCTGTGGGCTGC					
rat	GTTGGGTGCCCTGTGGCA--GGCTTGTCGAATGCCACTCTGTCCCCTCTCTCTGGCCCTAGGCCTGC					
	170	180	190	200	210	220
	280	290	300	310	320	330
human	GGCTGGCTGGAACCTCTCAACCCAGTGATCCCAATACCTGCAGCTTCTGGGAAAGCTTCACTACCACCAC					
rat	GTCTGGCTGGAACACTCAACTCCAATGATCCCAATGTCTGTACCTTCTGGGAAAGCTTCAACCAGACCAC					
	240	250	260	270	280	290
	350	360	370	380	390	400
human	CAAGGAGTCCCCTCCGCCCCCTTCAGCCTGCTCCCTCAGAGCCCTGCGAGCGCCCTGGGAGGGCCCC					
rat	TAAGGAGTCCCACCTTCGCCCCCTTCAGCCTGCCCCAGCCGAGTCTGCGACAGGCCCTGGGAAGACCCC					
	310	320	330	340	350	360
	420	430	440	450	460	470
human	CATACTTGC-CCAGCCCAAAA---CT--CAGA---GGAAACTCCTGGCT-TCTAGGGATTCTTCTGC					
rat	CACACCTGCGCTCAGCCTACGGTTGTCTACCGGACTGTGTACCGTCAGGTGGTGAAGATGGACTCCCGCC					
	380	390	400	410	420	430
	480	490	500	510	520	530
human	ATGGTCTGTGTGGGGCTG-GAGTGCACTGGCGAGATC-GTAGTGCACTGCAACCTCAAACAGGGAATGC					
rat	CACGCCTG---CAGTGCTGTGGGGTTACTACGAGAGCAGTGGAGC-CTGTGTCC-CACTCTG---TGC					
	450	460	470	480	490	500
	550	560	570	580	590	600
human	GCTTTCTATGCGCCCTCAGCCCAGAGTGTGTGAGTGGTGCCCTTCCTCTG-GCCTCCCTGGCCACACTGT					
rat	CCAGG-AGTGTGTCCACGGT-----GCTGTGTG--GCTCCTAATCGGTGCCAGTGTGCACAGGCTGG					
	510	520	530	540	550	560

FIG.34A

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human 620 630 640 650 660 670 680
      GGTGGTGAAGACGGACCACCGCCAGCGCCTGCAGTGCCTGGCTTCTATGAGAGCAGGGGGTTCTGT
rat   CCGGGTGACGACTGT-----TCCAGTG--AG-TGTGCT-CC-TGGAA--TGTGGGGACCACAG----TGT
      570 580 590 600 610

human 690 700 710 720 730 740 750
      GTCCCGCTCTGTGCCCAGGAGTGTGTCCATGGCCGTTGTGTGGCACCCA--ATCAGTGCCAATGTGTGCC
rat   GACAGGCTCTG---CCTC---TGTGGCAACAGCAGTTCCTGTGATCCAGGAGTGGGGTGTGTTTTTGCC
      620 630 640 650 660 670 680

human 760 770 780 790 800 810
      AGGCTGGCGGGGCGACGACTGTTCCAGTGCCCCGAAGTGCCTTCAGCCCTGTACCCC--TGGCTACTATG
rat   CCTCTGGC-----CTGCAG--CC---CCCCGA-CTGCCTTCAGCCTTG--CCCGATGGCCACTATG
      690 700 710 720 730

human 820 830 840 850 860 870 880
      GCCCTGCCTGCCAGTTCGCTGCCAGTGCCATGGGGCACCTGCGATCCCAGACTGGAGCCTGCTTCTG
rat   GTCTGCCTGCCAGTTTGATTGCCATTGCTATGGGGCATCCTGTGACCCCCGGGATGGAGCCTGCTTCTG
      740 750 760 770 780 790 800

human 890 900 910 920 930 940 950
      CCCCAGAGAGAACTGGGCCAGCTGTGACGTGTCCTGTTCCAGGGCACTTCTGGCTTCTTCTGCCCC
rat   CCCCCAGGGAGAACAGGACCCAG-----GGCACTGATGGCTTCTTCTGCCCC
      810 820 830 840 850

human 960 970 980 990 1000 1010 1020
      AGCACCCATCCTTGCCAAAATGGAGGTGTTCCAAACCCACAGGGCTCCTGCAGCTGCCCCCTGGCT
rat   AGAATTATCCTTGCCAAAATGGAGGTGTTCTCAGGGCTCTCAAGGCTCCTGCAGCTGCCACCGGGCT
      860 870 880 890 900 910 920

human 1030 1040 1050 1060 1070 1080 1090
      GGATGGGCACCATCTGCTCCCTGCCCTGCCAGAGGGCTTTCACGGACCCAACTGCTCCAGGAATGTGCG
rat   GGATGGGTGTCATCTGTTCCCTGCCATGCCAGAGGGTTTCACGGACCCAACTGTACTCAGGAATGTGCG
      930 940 950 960 970 980 990

human 1100 1110 1120 1130 1140 1150 1160
      CTGCCACAACGGCGGCCTCTGTGACCGATTCACTGGGCAGTGCCGCTGCGCTCCGGGTACACTGGGGAT
rat   TTGCCACAATGGTGGCCTTTGTGACAGGTTTACTGGGCAGTGCCACTGTGCTCCTGGCTATATCGGGGAT
      1000 1010 1020 1030 1040 1050 1060

human 1170 1180 1190 1200 1210 1220 1230
      CGGTGCCGGGAGGAGTGCCCGGTGGGCGCTTTGGGCAGGACTGTGCTGAGACGTGCGACTGCGCCCCGG
rat   CGGTGCCGTGAAGAGTGCCCTGTGGGCGCTTCGGTCAAGACTGTGCTGAGACCTGTGACTGTGCTCCTG
      1070 1080 1090 1100 1110 1120 1130

human 1240 1250 1260 1270 1280 1290 1300
      ACGCCCGTTGCTTCCCGGCCAACGGCGCATGTCTGTGCGAACACGGGTTCACTGGGGACCGCTGCACGGA
rat   GCGCTCGTTGCTTTCCCTGCCAATGGCGCGTGTCTGTGCGAACATGGCTTCACAGGCGACCGCTGCCTGA
      1140 1150 1160 1170 1180 1190 1200

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FIG.34B

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      1310      1320      1330      1340      1350      1360      1370
human  TCGCCTCTGCCCCGACGGCTTCTACGGTCTCAGCTGCCAGGCCCCCTGCACCTGCGACCGGGAGCACAGC
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    GCGACTCTGTCCAGATGGCCGCTATGGTCTGAGCTGCCAAGATCCCTGCACCTGCGACCCAGAACACAGT
      1210      1220      1230      1240      1250      1260      1270

      1380      1390      1400      1410      1420      1430      1440
human  CTCAGCTGCCACCCGATGAACGGGGAGTGCTCCTGCCTGCCGGGCTGGGCGGGCCTCCACTGCAACGAGA
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    CTCAGCTGCCACCCAATGCACGGCGAGTGCTCCTGCCAGCCAGGTTGGGCGGGCCTCCACTGCAACGAGA
      1280      1290      1300      1310      1320      1330      1340

      1450      1460      1470      1480      1490      1500      1510
human  GCTGCCCGCAGGACACGCATGGGCCAGGGTGCCAGGAGCACTGTCTCTGCCTGCACGGTGGCGTCTGCCA
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    GCTGCCCTCAGGACACGCACGGAGCCGGTTGCCAGGAGCACTGCCTCTGTCTGCACGGCGGTGTTTGCCT
      1350      1360      1370      1380      1390      1400      1410

      1520      1530      1540      1550      1560      1570      1580
human  GGCTACCAGCGGCCTCTGTCACTGCGCGCGGGTTACACGGGCCCTCACTGTGCTAGTCTTTGTCTCCT
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    CGCCGACAGCGGCCTCTGCCGGTGTGCACCTGGCTACACGGGACCTCACTGCGCTAATCTTTGTCCACCT
      1420      1430      1440      1450      1460      1470      1480

      1590      1600      1610      1620      1630      1640      1650
human  GACACCTACGGTGTCAACTGTTCTGCACGCTGCTCATGTGAAATGCCATCGCTGCTACCCATCGAGC
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    AACACTTATGGGATCAACTGTTCTCCCACTGCTCCTGTGAAATGCCATTGCTGCTCTCCTGTCCGACG
      1490      1500      1510      1520      1530      1540      1550

      1660      1670      1680      1690      1700      1710      1720
human  GCGAGTGCGTCTGCAAGGAAGGTTGGCAGCGTGGTAACTGCTCTGTGCCCTGCCACCCGGAACCTGGGG
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    GCACGTGCATCTGCAAGGAAGGTTGGCAGCGTGGTAACTGCTCTGTGCCCTGTCCCCCTGGCACCTGGGG
      1560      1570      1580      1590      1600      1610      1620

      1730      1740      1750      1760      1770      1780      1790
human  CTTCACTTGCAATGCCAGCTGCCAGTGTGCCCATGAGGCAGTCTGCAGCCCCAACTGGAGCCTGTACC
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    CTTCACTTGCAATGCCAGTGTGCCAGTGTGCCACGAGGGAGTCTGCAGCCCCAACTGGAGCCTGTACT
      1630      1640      1650      1660      1670      1680      1690

      1800      1810      1820      1830      1840      1850      1860
human  TGCACCCCTGGGTGGCATGGGGCCCACTGCCAGCTGCCCTGTCCGAAGGGGAGTTTGGAGAAGGTTGTG
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    TGCACCCCTGGGTGGCGTGGGGTTCACTGCCAACTTCCGTGCCCGAAGGGACACTTTGGTGAAGGTTGTG
      1700      1710      1720      1730      1740      1750      1760

      1870      1880      1890      1900      1910      1920      1930
human  CCAGTCGCTGTGACTGTGACCACTCTGATGGCTGTGACCCTGTTTCATGGACGCTGTCACTGCCAGGCTGG
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    CCAGTGTCTGTGACTGTGACCACTCCGATGGCTGTGACCCTGTTTCATGGACACTGCCGATGTCAAGGCTGG
      1770      1780      1790      1800      1810      1820      1830

      1940      1950      1960      1970      1980      1990      2000
human  CTGGATGGGTGCCCGCTGCCACCTGTCCTGCCCTGAGGGCTTATGGGAGTCAACTGTAGCAACACCTGC
      ::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::::
rat    CTGGATGGGCACACGTTGCCACCTGCCTTGCCAGAGGGCTTTTGGGAGCCAACTGCAGCAATGCCTGT
      1840      1850      1860      1870      1880      1890      1900
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FIG.34C

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human 2010 2020 2030 2040 2050 2060 2070
      ACCTGCAAGAATGGGGGCACCTGTCTCCCTGAGAATGGCAACTGCGTGTGTGCACCCGGATTCCGGGGCC
      .....
rat    ACCTGCAAGAATGGTGGCACTTGTGTACCTGAGAACGGCAACTGTGTGTGCGCACCAGGGTTTCAGAGGCC
      1910 1920 1930 1940 1950 1960 1970

human 2080 2090 2100 2110 2120 2130 2140
      CCTCCTGCCAGAGATCCTGTGCAGCCTGGCCGCTATGGCAAACGCTGTGTGCCCTGCAAGTGCCTAACCA
      .....
rat    CCTCCTGCCAGAGGCCCTGCCCGCCTGGTCCGTATGGCAAACGCTGTGTGCCCTGCAAGTGCACCAACCA
      1980 1990 2000 2010 2020 2030 2040

human 2150 2160 2170 2180 2190 2200 2210
      CTCCTTCTGCCACCCCTCGAACGGGACCTGCTACTGCCTGGCTGGCTGGACAGGCCCGACTGCTCCCAG
      .....
rat    TTCTTCTGCCACCCCTCGGATGGGACCTGCTCCTGCCTGGCAGGCTGGACAGGCCCTGACTGCTCTGAA
      2050 2060 2070 2080 2090 2100 2110

human 2220 2230 2240 2250 2260 2270 2280
      CCATGCCCTCCAGGACACTGGGGAGAAAACCTGTGCCAGACCTGCCAATGTCAACATGGTGGGACCTGCC
      .....
rat    CATGTCCCCCAGGCCACTGGGGACTCAAATGCTCCCAACCCTGCCAGTGTATCATGGTGCCACCTGCC
      2120 2130 2140 2150 2160 2170 2180

human 2290 2300 2310 2320 2330 2340 2350
      ATCCCCAGGATGGGAGCTGTATCTGCCCCCTAGGCTGGACTGGACACCACTGCTTAGAAGGCTGCCCTCT
      .....
rat    ACCCCCAGGATGGGAGCTGTGTCTGCATCCAGGCTGGACTGGACCCAACCTGCTCGGAAGGCTGCCCATC
      2190 2200 2210 2220 2230 2240 2250

human 2360 2370 2380 2390 2400 2410 2420
      GGGGACATTTGGTGCTAACTGCTCCCAGCCATGCCAGTGTGGTCTGGAGAAAAGTGCCACCCAGAGACT
      .....
rat    AAGAATGTTTGGTGTCAACTGCTCCCAGCTATGTCAAGTGTATCTGGAGAGATGTGCCACCCAGAGACT
      2260 2270 2280 2290 2300 2310 2320

human 2430 2440 2450 2460 2470 2480 2490
      GGGGCCTGTGTATGTCCCCCAGGGCACAGTGGTGCACCTTGCAGGATTGGAATCCAGGAGCCCTTTACTG
      .....
rat    GGGGCTTGCGTCTGTCCCCCAGGACACAGTGGTGGCACTGCAAAGTGGGCAGCCAGGAGTCTTCACCA
      2330 2340 2350 2360 2370 2380 2390

human 2500 2510 2520 2530 2540 2550 2560
      TGATGCCGACCACTCCAGTAGCGTATAACTCGCTGGGTGCAGTGATTGGCATTGCAGTGCTGGGGTCCCT
      .....
rat    TAATGCCACCTCTCCTGTGATCCATAACTCACTGGGTGCCGTGATTGGCATTGCAGTGCTGGGGACCTT
      2400 2410 2420 2430 2440 2450 2460

human 2570 2580 2590 2600 2610 2620 2630
      TGTGGTAGCCCTGGTGGCACTGTTTCATTGGCTATCGGCACTGGCAAAAAGGCAAGGAGCACCACCACCTG
      .....
rat    TGTGGTGGCCCTGGTAGCACTGTTTATTGGCTACCGCACTGGCAAAAAGGCAAGGAACATGAGCACTTG
      2470 2480 2490 2500 2510 2520 2530

human 2640 2650 2660 2670 2680 2690 2700
      GCTGTGGCTTACAGCAGCGGGCGCCTGGACGGCTCCGAGTATGTCATGCCAGATGTCCCTCCGAGCTACA
      .....
rat    GCAGTGGCTTACAGCACTGGGCGACTGGATGGCTCCGATTACGTCATGCCAGATGTCTCTCCGAGCTACA
      2540 2550 2560 2570 2580 2590 2600

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FIG. 34D

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      2710      2720      2730      2740      2750      2760      2770
human  GTCACTACTACTCCAACCCAGCTACACACCCTGTGCGAGTGTCTCCCAACCCCAACCCCTAACAA
      .....
rat    GTCACTACTATTCCAACCCAGCTACACACACTGTCTCAGTGTCTCTAACCCTCCACCCCTAACAA
      2610      2620      2630      2640      2650      2660      2670

      2780      2790      2800      2810      2820      2830      2840
human  GGTTCAGGC---CCGCTCTTTGCCAGCCTGCAGAACCTGAGCGGCCAGGTGGGGCCCAAGGGCATGAT
      .....
rat    GATTCCAGGCAGTCAGCTGTTTGTCACTCCAGGCATCTGAGCGGCCAAACAGAAACCATGGGCGAGAT
      2680      2690      2700      2710      2720      2730      2740

      2850      2860      2870      2880      2890      2900      2910
human  AACCACACCACCTGCCTGCTGACTGGAAGCACCGCCGGGAGCCCCCT-CCAGGGCCrCTGGACAGGGGG
      .....
rat    AACCACGCCACACTGCCCGCTGACTGGAAGCACCGAGCGGAGTCCCATGACAGAGC--TTTCTCAGGC
      2750      2760      2770      2780      2790      2800

      2920      2930      2940      2950      2960      2970
human  AGCAGCCGCCTGGACCGAAG-----CTACAGCTATAGCTACAGCAATGGCCAGGCCATTCTACGATA
      .....
rat    ACCAGCCACCTGGACCGAAGGTATAGCTGTAGCTATGGCCACAGGAATGGCCCGGGGCCATTCTGTCATA
      2810      2820      2830      2840      2850      2860      2870

      2980      2990      3000      3010      3020      3030      3040
human  AAGGGCTCATCTCTGAAGAGGAGCTCGGGGCCAGTGTGGCTTCCCTGAGCAGTGAGAACCCATATGCCAC
      .....
rat    AAGGTCCCATGTGTGAAGAAGGACTAGGGGCAAGCGTTATGTCCCTGAGCAGTGAGAACCCCTATGCGAC
      2880      2890      2900      2910      2920      2930      2940

      3050      3060      3070      3080      3090      3100      3110
human  CATCCGGGACCTGCCAGCTTGCCAGGGGGCCCCCGGAGAGCAGCTACATGGAGATGAAAGGCCCTCCC
      .....
rat    CATCCGAGACCTGCCCGCTGCCGGGAAACCCGAGAAAGCAGCTATGTGGAGATGAAAGGCCCTCCA
      2950      2960      2970      2980      2990      3000      3010

      3120      3130      3140      3150      3160      3170      3180
human  TCAGGATCTGCCCCAGGCAGCCTCCTCAGTTTTGGGACAGCCAGAGCGGGCGCAACCCAGCCACAGA
      .....
rat    TCAGTGTCTCCCCCAGGCAGCCTCTTCATCTCCGGGACAGGCAG--CAGCAGCAACTGCAGTCTCAGA
      3020      3030      3040      3050      3060      3070      3080

      3190      3200      3210      3220      3230      3240      3250
human  GAGACAGTGGCACCTACGAGCAGCCAGCCCCCTGATCCATGACCGAGACTCTGTGGCTCCCAGCCCC
      .....
rat    GAGACAGCGGCACCTATGAGCAGCCCACTCCCTTGAGCCGTAATGAAGAGTCTGTGGGCTCCATGCCCC
      3090      3100      3110      3120      3130      3140      3150

      3260      3270      3280      3290      3300      3310      3320
human  TCTGCCTCCGGGCCTACCCCCGGCCACTATGACTCACCAAGAACAGCCACATCCCTGGACATTATGAC
      .....
rat    TCTTCCTCCGGGCCTGCCACCCGGCCACTATGACTCGCCCAAAACAGCCACATCCCTGGACACTATGAC
      3160      3170      3180      3190      3200      3210      3220

      3330      3340      3350      3360      3370      3380      3390
human  TTGCCTCCAGTACGGCATCCCCATCACCTCCACTTCGAGCGCAGGACCGTTGAGGAGCCAGGATGGTAT
      .....
rat    TTGCCTCCAGTACGGCATGGTGGATCACCTCCATCCCGGCGCAGGACCGCTGAGGAGCCAGCATGGTAT
      3230      3240      3250      3260      3270      3280      3290

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FIG.34E

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human 3400 3410 3420 3430 3440 3450 3460
      GGCAGAGGCCAGCACACCTGGCTGTTGCTGCTCAAGGCTGGGGACAGAGCCTAGTGTACCCCTGCCAGGA
rat   GG--GAG-----AGTGCCT-GTGAACCC-TGCCAGGA
      3300 3310 3320

human 3470 3480 3490 3500 3510 3520 3530
      GCAGGGAGTGGACCGGCAGGCTGTGAACATGAACAACGCTTAACAGAGCAAGTGATGGGAGCCTTGTTC
rat   GCAGGGCCTGGACCGCAGGC-----CATGAA-----TAGACATA-----
      3330 3340 3350

human 3540 3550 3560 3570 3580 3590 3600
      TGGGTTCTACCATGGGAGACGCTGATCAGCAGGATGCCTGGCTCCCTTCCCAACCCACTGCTCCCAAGG
rat   -----CTTGG-----TGAA-----
      3360

human 3610 3620 3630 3640 3650 3660 3670
      CCTCCAGGGCCCTGTGTACATAAACTGGTGGGTTGGAAGTTGCTGGGTAACCTGATTTTCAGACATGCGT
rat   -----GTGAACCGAGACTG-AGGATGG-----
      3370 3380

human 3680 3690 3700 3710 3720 3730 3740
      GTGGGGTACCTTTTCTGTGCATGCTCAGCCTGGGCTCTGTGCGTGTGTGTTTCTGTGATTTTAGAAGG
rat   -----CTCTGC-----
      3390

human 3750 3760 3770 3780 3790 3800 3810
      GTACCAGGCAGGTTCTGTCTAGGGCACTTACCATTAGTAGGGAGATGGAACCAACCCAATTAACCTCTA
rat   -TTCCA-----CCGAGGG-----AGACACTA
      3400 3410

human 3820 3830 3840 3850 3860 3870 3880
      GCAATAGCCTCCTAACTGGCCTCCTCATTGATTAGTGAACCTTCCAATGCATGGCTCATAATTTCAA
rat   G-----TTGGC-----
      3420

human 3890 3900 3910 3920 3930 3940 3950
      ATACAGGCTGGTTAGTTACTCCCTACCTGAAAGCCTTCATAGGTGCCTCTTTGCTCTTCTGCCAGTATCA
rat   -----AAAG-----

human 3960 3970 3980 3990 4000 4010 4020
      AAACCTTTTGAAGGCCCTTAAGGCCCTGCTTTGCCTGGCCCATCTGTCTCTCCAGCCTCACCTTGAACGTG
rat   -----TGTCT-----
      3430

human 4030 4040 4050 4060 4070 4080 4090
      GTTCCTGTCACTGCACGCCAGTCACACCGGCCCTCTAGTCTCTGTAGGCCACTCTTCTTTCTGGCACA
rat   -----AACCTCC-----

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FIG.34F

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      4100      4110      4120      4130      4140      4150      4160
human  GGGACCTGCACACCTGGAGTGCCCTTCCTCCCCCACTCGCCTGTTACACCCCTGCTTTTCTTTACACCTC
rat    -----CTTTTCC-----
                               3440

      4170      4180      4190      4200      4210      4220      4230
human  CTCCTCAGGGAAGTGCCACCCCTCCGTACATCTTTCACAGCCCTGATTGCAGCTGTGTTCACTCACCAGG
rat    -----AGCCC-ATTGCT-----CAAG
                               3450

      4240      4250      4260      4270      4280      4290      4300
human  TACCTGCAGAAGGCCTACAGGGTGCCAGGCACCTCTTTAATGGGTTCTTTCTTTATGTGATTATTTGATT
rat    T-----
      3460

      4310      4320      4330      4340      4350      4360      4370
human  AATCTCTGCCTCCCCACTAGACTGTAAGCTCCCTGAAGGCAAGAATCCTGTGCTTATGCTCAATATTAG
rat    -----CCCCA-----

      4380      4390      4400      4410      4420      4430      4440
human  CTCTCCCTTGGCACAGAGTAGGCACTCAACAAATGCTCCCCAAAAGGCTGAGTGGCTGACTGAATTAAGT
rat    -----GGCTGTG-----
                               3470

      4450      4460      4470      4480      4490      4500      4510
human  ACCAGTGACATGCAGTAACTGCTAAGATAGATGAGCCATCTGTATGCTCTGACAGTTACAGACTGAATAA
rat    -----GACATG-----

      4520      4530      4540      4550      4560      4570      4580
human  GTTGGAGACTTCCCTAAAGGGTGGCATTTCGCCAGGGTAACAACGCAGAGCTCAGGTGTGGGAAGGTGCC
rat    -----

      4590      4600      4610      4620      4630      4640      4650
human  AGGGGCAGGGGTGCAGAGGGGCTGAGGCTGAGGGGGTGCAGAGGCTGGAGAAAGGATAACAGGAGAGAG
rat    -----AGCTGGTGG-----
                               3480

      4660      4670      4680      4690      4700      4710      4720
human  TATACAGGCATGCCCTTGATTTATTGCACCTCACAGGTAGCAGAAATTTTAAAGAAATTGAAGGTTTGGG
rat    -----GCAGAAATGTT-----GTTGTTGAAG-----
                               3490      3500

      4730      4740      4750      4760      4770      4780      4790
human  ACATATATGTGACAGCAATAGGTTAAGAAAAGCAAAGCAGAGAAATTGAAGATTTGTGTCAACACTGCTT
rat    -----

```

FIG.34G

```

      4800      4810      4820      4830      4840      4850      4860
human TAAGCAAATCTGTTGGCACCATTTCCTCAATAGCATGTGCCCATTTGGGTCTCTACATTGCATTTGGT
rat   .....TCTG.....ATTTAGAT.....
           3510                       3520

      4870      4880      4890      4900      4910      4920      4930
human AATTGCTTGCAATATTTCAAGCATTTTCATTGTTATTATATGTGTTATAGTGATCTGTGATCAGTGATCT
rat   .....

      4940      4950      4960      4970      4980      4990      5000
human TTGATATATTATTGTAATTGTTTCGGGGCGCCATGAACCGCACCCATATAACACGGTAAACTTAATCAGC
rat   -TGATTTTTTAAAAAAA-.....
           3530

      5010      5020      5030
human AAAAAAAAAAAAAAAAAAAGGGCGGCCG-
rat   AAAAAAAAAAAAAAAAAAAGGGCGGCCGC
      3540      3550      3560
```

FIG.34H

inputs GTC-GACCCACGCGTCCGG---TGACCCTGTTCATGGACAGT-----GCCGATGTCAGG--CTGGT---

10 20 30 40 50 60 70

inputs TGGATGGGCACA-CGCTGCCAC--CTGCCTTG-CCCGBA--GG-GCTTTTGGGGAG-CCAAC-TGCAG

80 90 100 110 120 130 140

inputs -TAACACCTGTACC-TGCAAGAATGGTGGTACCTGTG--TGTCT-GAGAATGGCAACTGCCTGTGCGCAC

150 160 170 180 190 200

inputs CAG----GGTCCGAGGCCCC-CTCCTGCCAGAGGCCCTGCCCGCC--TGGTCGCTATGGCAA-AC--GCT

210 220 230 240 250 260 270

inputs GTGT--GCAATGC-----AAGTGT--AACAAACAACCATCTCTCTGCCACCCATCG-----

280 290 300 310 320 330 340

inputs -GACGGGACCTG-----CTCCT-GCCTG--GCGGGCTG-GACAGGC--CCTGACTGC--TCCG--AG

350 360 370 380 390 400 410

inputs GC-----ATG--TCCC--CCAGGCCA-----CTGGGG-----ACT-CAAATGCT-----CC----

420 430 440 450 460 470 480

inputs --CAACTCTG--CCAG-----TGTCATCA-----TG-GTGGGACCT-----GCCA-----CCCC--

490 500 510 520 530 540 550

TCCCACTCTGTGCCAGGAGTGTGTCCACGGTCGCTGTGTGGCTCCTAATCGGTGCCAGTGTGTGCACAGG

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      420      430      440      450      460
inputs CAGGATGGGAG---CTGTATC-----TGCACGCCAGGCTGGACTGGACC-CAA-----CTGC
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      CTGGCGGGGTGACGACTGTTCCAGTGAGTGTGCTCCTGGAATGTGGGACCACAGTGTGACAGGCTCTGC
      560      570      580      590      600      610      620

      470      480      490      500
inputs TTGGAAGGCTGC-----CCA-----CCAAGAATGTTTGGTGT-----CAACTGCTCC
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      CTCTGTGGCAACAGCAGTTCCTGTGATCCAGGAGTGGGGTGTGTTTTTGGCCCTCTGGCCTGCAGCCCC
      630      640      650      660      670      680      690

      510      520      530
inputs C-----AGCTATGTC--AGTG-----TGATCT-----CGGAGAGATG-----TGC-----
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      CCGACTGCCTTCAGCCTTGCCCCGATGGCCACTATGGTCTGCTGCCAGTTTGATTGCCATTGCTATGG
      700      710      720      730      740      750      760

      540      550      560      570      580
inputs --CACCCAGAGAC-----TGGGGCTTGTCTGTCCCCCAGG-----ACACAG-----TGGTG
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      GGCATCCTGTGACCCCCGGGATGGAGCCTGCTTCTGCCCCCAGGGAGAACAGGACCCAGGGCACTGATG
      770      780      790      800      810      820      830

      590      600      610      620
inputs -----CAGAC-----TGCAAAATGGGAAG---CC--AGGAGTC-CTT--CACCATAA-
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      GCTTCTTCTGCCCCAGAACTTATCCTTGCCAAATGGAGGTGTTCTCAGGGCTCTCAAGGCTCCTGCAG
      840      850      860      870      880      890      900

      630      640      650
inputs -TGCCACC-----TCT---CCCG---TGACCCATAA-----CTC-----ACTGG
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      CTGCCCACCGGGCTGGATGGGTGTCATCTGTTCCCTGCCATGCCAGAGGGTTTCCACGGACCCAACTGT
      910      920      930      940      950      960      970

      660      670      680      690      700      710
inputs GTGCAGTGATTGGCATTGCAGTACTGGGAACCTCGTG---GTGGCCCTGATAG---CACTGTTTCA-T
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      ACTCAG-GAATGTCGTTGCCACAATGGTGGCCTTTGTGACAGGTTTACTGGGCAGTGCCACTGTGCTCCT
      980      990      1000      1010      1020      1030      1040

      720      730      740
inputs GGCTA-----CCG-----CCAGTGG-----CAAAA--GGGCAAGGAACA
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      GGCTATATCGGGATCGGTGCCGTGAAGAGTGCCCTGTGGGCCGCTTCGGTCAAGACTGTGCTGAGACCT
      1050      1060      1070      1080      1090      1100      1110

      750      760      770      780      790
inputs ----TGAGCACTTGGCA--GTGGCTTAC-----AGCACTGGGCGG--CTGG-ATGGCTCTGATTA
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
      GTGACTGTGCTCCTGGCGCTCGTTGCTTTCTGCCAATGGCGCGTGTCTGTGCGAACATGGCTTCACAGG
      1120      1130      1140      1150      1160      1170      1180

```

FIG.35B

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```

      800      810      820      830      840      850
inputs  CGTCA--TGC-CAGAT-GTCTCT--CCGA-----GCTATAGTCACTACTACT-----CCAACCCCAGC
      1190      1200      1210      1220      1230      1240      1250
      CGACCGCTGCACTGAGCGACTCTGTCCAGATGGCCGCTATGGTCTGAGCTGCCAAGATCCCTGCACCTGC
      1190      1200      1210      1220      1230      1240      1250

      860      870      880      890      900
inputs  TACC--ACACACTGTCTCAGTGTTCCTTAACCGCCCGC----CCCCTAACA--AGGTCC--CAGGCA
      1260      1270      1280      1290      1300      1310      1320
      GACCCAGAACACAGTCTCAGCTGCCACCCAATGCACGGCGAGTGCTCCTGCCAGCCAGGTTGGGCGGGCC
      1260      1270      1280      1290      1300      1310      1320

      910      920      930      940      950
inputs  G--TCAGCT-CTTTGTCACTCTCAGGCC-C---CTGAGC---GGCCA--AGCAGAGCC-----CA
      1330      1340      1350      1360      1370      1380      1390
      TCCACTGCAACGAGAGCTGCCCTCAGGACACGCACGGAGCCGGTTGCCAGGAGCACTGCCTCTGTCTGCA
      1330      1340      1350      1360      1370      1380      1390

      960      970      980      990      1000      1010
inputs  CGGGCGTCAGAACCATACCACACTGC--CCGCTGACTGGAAGCACC--GC---CGGGAGCCC-----C
      1400      1410      1420      1430      1440      1450      1460
      CGGCGGTGTTTGCCTCGCCG-ACAGCGGCCTCTGCCGGTGTGCACCTGGCTACACGGGACCTCACTGCGC
      1400      1410      1420      1430      1440      1450      1460

      1020      1030      1040      1050      1060
inputs  ATGACAGAGGC-GCCAGCCAC-----CTGGACCGAA-GCTATAGCTGTA----GCTATAGCC
      1470      1480      1490      1500      1510      1520      1530
      TAATCTTTGTCCACCTAACACTTATGGGATCAACTGTTCTCCCACTGCTCCTGTGAAATGCCATTGCC
      1470      1480      1490      1500      1510      1520      1530

      1070      1080      1090      1100      1110
inputs  A-----CAGG-AATGGCCCAGG--AC--CATT-----CTGTCATAAAGGTCCCATCTCTGAA---GA-
      1540      1550      1560      1570      1580      1590      1600
      TGCTCTCCTGTGACGGCACGTGCATCTGCAAGGAAGTTGGCAGCGTGGTAACTGCTCTGTGCCCTGTC
      1540      1550      1560      1570      1580      1590      1600

      1120      1130      1140      1150      1160
inputs  -----GGGACTAGGGGCAAGCGTTA-TGTCCCTGA-GCAGTGAGAACCC-CTA-----TGCTACC---
      1610      1620      1630      1640      1650      1660      1670
      CCCCTGGCACCTGGGGCTTCAGTTGCAATGCCAGTTGCCA-GTGTGCCACGAGGGAGTCTGCAGCCCCC
      1610      1620      1630      1640      1650      1660      1670

      1170      1180      1190      1200      1210
inputs  -ATCCGAGACCTG-----CCCAGCCTGCC-TGGGGAAC---CC-----CGAG--AAAGTGGCT
      1680      1690      1700      1710      1720      1730      1740
      AAAGTGGAGCCTGTACTTGCACCCCTGGGTGGCGTGGGTTCACTGCCAACTTCCGTGCCGAAGGGACA
      1680      1690      1700      1710      1720      1730      1740

      1220      1230      1240      1250      1260
inputs  ATGTGGAGATGAAAGGACC---TCCAT--CAGTGTCCCCTCCA-GGCAGT---CTCTTCAT-----C
      1750      1760      1770      1780      1790      1800      1810
      GTTTGGTGAAGGTTGTGCCAGTGTCTGTGACTGTGACCACTCCGATGGCTGTGACCCTGTTTCATGGACAC
      1750      1760      1770      1780      1790      1800      1810

```

FIG.35C

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```

      1270      1280      1290      1300      1310
inputs T-CCGG-GACAGGCAG-CAG-----CGG---CAACTGC--AGCCACAGAGGG--ACAGCGGCACC
      1820      1830      1840      1850      1860      1870      1880
      TGGCGATGTCAGGCTGGCTGGATGGGCACACGTTGCCACCTGCCTTGCC--CAGAGGGCTTTTGGGGAGCC

      1320      1330      1340      1350
inputs TA-TG-AGCA--GCC-----CAGC-----CCCTTGAG--CCATAATGAAGAGTCTTTGGG----
      1890      1900      1910      1920      1930      1940      1950
      AACTGCAGCAATGCCTGTACCTGCAAGAATGGTGGCACTTGTGTACCTGAGAACGGCAACTGTGTGTGCG

      1360      1370      1380      1390      1400
inputs CTCCA-----C---GCCCCGCTTCCTCCAGGCCTGCC--TCCTGGTCACTACGACT--C-----CC
      1960      1970      1980      1990      2000      2010      2020
      CACCAGGGTTTCAAGAGGCCCTCCTGCCAGAGGCCCTGCCCGCCTGGTGGCTATGGCAAACGCTGTGTGCC

      1410      1420      1430      1440      1450
inputs C--CAAG---AACAGCCATA-TCCCTG-----GAC-----ACTATGACTTGCCT--C---CAGTAC-
      2030      2040      2050      2060      2070      2080      2090
      CTGCAAGTGCAACAACCAATTCTTCTGCCACCCGTCGGATGGGACCTGCTCCTGCTGGCAGGCTGGACA

      1460      1470      1480
inputs GGC---ATC--CTC-----CAT---CCCCT--CCA-----TCCCGGC---GCCAG-GAC
      2100      2110      2120      2130      2140      2150      2160
      GGCCCTGACTGCTCTGAATCATGTCCCCAGGCCACTGGGGACTCAAATGCTCCCAACCCTGCCAGTGTG

      1490      1500      1510      1520      1530      1540
inputs CGC-TGAAGA-GCCGGCAT-----GGTATGGGAGC-GTGCCTATGTACCTTGC---CAGGA-----G
      2170      2180      2190      2200      2210      2220      2230
      ATCATGGTGCCACCTGCCACCCCAAGGATGGGAGCTGTGTCTGCATCCAGGCTGGACTGGACCCAAGTG

      1550      1560      1570      1580
inputs CAGGGACTG--GACCAGCAGG-----CCACG-----AACAGAAACA-----CTTGGTGAA
      2240      2250      2260      2270      2280      2290      2300
      CTCGGAAGGCTGCCCATGAAGAATGTTTGGTGTCAACTGCTCCCACTATGTCAAGTGTATCTTGGAGAG

      1590 1600 1610 1620 1630
inputs GTGAAC-----AGAGACGGACTGTGGC-CCTGTGCTTC---CACCGAGGGAGACACT---AGTTGACA
      2310      2320      2330      2340      2350      2360      2370
      ATGTGCCACCCAGAGACTGGGGCTTGGCTGTGTCCCCAGGACACAGTGGTGGCCTGCAAAAGTGGGCA

      1640      1650      1660      1670      1680      1690
inputs ---AAGTGTCTAAC-CCTCTTTTCCAACC-CAC---TGCTC---AAGTCCCTGTGGAC---ATAAGC--
      2380      2390      2400      2410      2420      2430      2440
      GCCAGGAGTCTTCAACATAATGCCACCTCTCCTGTGATCCATAACTCACTGGGTGCCGTGATTGGCAT

```

FIG.35D

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```

      1700      1710      1720      1730      1740
inputs TGGTGGGCAGAA-----TGTTGTTGTACAAGTG---TGATTTTAG---ATCGATTTTTTTTTAAAGT-
      2450      2460      2470      2480      2490      2500      2510
      TGCAGTGTCTGGGGACCCCTTGTGGTGGCCCTGGTAGCACTGTTTATTGGCTACCGACACTGGCAAAGGGC

      1750      1760      1770      1780      1790      1800      1810
inputs ATGTGTTGGGTAC-CTTTTCTGTG-TGTATGCTCAGGCAGGCTGTGTGTCTCTAGTTGGCTTTAGAG
      2520      2530      2540      2550      2560      2570      2580
      AAGGAACATGAGCACTTGGCAGTGGCTACAGCACTGGGCGACTGGATGGCTC-CGATTACGTCATGCCA

      1820      1830      1840      1850      1860      1870
inputs GGAGTC-----AGGTATAGGTTCTGCCTT--CTGCACT---TTCCA-TCT-TATCT-AGTAGTCAGCTT
      2590      2600      2610      2620      2630      2640      2650
      GATGTCTCTCCGAGCTACAGTCACTACTATTCCAACCCTAGCTACCAACACTGTCTCAGTGTCTCTCTA

      1880      1890      1900      1910      1920
inputs -CCAAGCTTAAGTAGTTAGAGCTCCA--C---CAGCAG-----CAG-GCCCTAACTAC---CTGCCTGC
      2660      2670      2680      2690      2700      2710      2720
      ACCCTCCACCCCTAACAAGATTCCAGGCAGTCAGCTGTTTGTGAGCTCCAGGCATCTGAGCGGCCAAA

      1930      1940      1950      1960      1970
inputs CCTTCACC-----C-AGTAATCCTC-CATGTCTTTGCTCAGA-GGATTGCTCC-CCGA----CTCT----
      2730      2740      2750      2760      2770      2780      2790
      CAGAAACCATGGGCGAGATAACCCAGCCACACTGCCCGCTGACTGGAAGCACCGAGGGAGTCCCATGAC

      1980      1990      2000      2010      2020
inputs GGTGTTGTCCTCTG---GTACGCCTTGAC---GGTCCTGCAGT--CT---CC-C-----TTTCCCG
      2800      2810      2820      2830      2840      2850      2860
      AGAGCTTTCTCAGGCACCAAGCCACCTGGACCGAAGGTATAGCTGTAGCTATGGCCACAGGAATGGCCCG

      2030      2040      2050      2060      2070      2080
inputs T---CTTGCT-TCATT-----CTTTCCAGAAATGAAGGCTGTCTGCCACCCTACT-TCCCAGCCCAGGA
      2870      2880      2890      2900      2910      2920      2930
      GGGCCATTCTGTCTATAAAGGTCCCATCTCTGAAGAAGGACTAGGGGCAAGCGTTATGTCCCTGAGCAGTG

      2090      2100      2110      2120      2130      2140
inputs A-----TTGGCA--CATCTAAGTTCAGCC-----TTCCTAAGTTACCGTTGAGTCCTGCTTGCCCTT
      2940      2950      2960      2970      2980      2990      3000
      AGAACCCCTATGCGACCATCCGAGACCTGCCCGGCTGCCTGGGGAACCCGAGAAAGCAGCTATGTGGA

      2150      2160      2170      2180      2190      2200
inputs CACATAT-----TCCA-CAGAA-CACCCACC-----CCACATCTGCTTCATAGCTACTCTCTTCTCCAC
      3010      3020      3030      3040      3050      3060      3070
      GATGAAAGGCCCTCCATCAGTGTCTCCCCCAGGCAGCCTCTTCATCTCCGGGACAGGCAGCAGCAGCAA

```

FIG.35E

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```

      2210      2220      2230      2240      2250      2260
inputs  GTACCCACAGAAGGCAGAAGTGGTACCAGGCAAGAAGATGGGA---TTGTTGCATTTTGTGTTTTG
      3080      3090      3100      3110      3120      3130      3140
      CTGCAGTCTCAGAGAGACAGCGGCACCTAT-GAGCAGCCCACTCCCTTGAGCCGTAATGAAGAGTCTGTG
      2270      2280      2290      2300      2310      2320      2330
inputs  AGACTCTGT-CTCACTATGTAGTCCTGGCTGGCCTG--GAACTCAAGAGCTCTGCCTGCCTCTGCCTCTT
      3150      3160      3170      3180      3190      3200      3210
      GG-CTCCATGCCCCCTCT-TCCTCCGGGCCTGCCACCGGCCACTATGACTCGCCAAAAACAGCCACAT
      2340      2350      2360      2370      2380
inputs  ----GAGTGCTGGGTTA-----ACGGCT--CAGGGTCACATGCA---CAGCTCAAGCTGCACT--
      3220      3230      3240      3250      3260      3270      3280
      CCCTGGACACTATGACTTGCCTCCAGTACGGCATCCTCCATCACCTCCATCCCGGCCAGGACCGCTGA
      2390      2400      2410      2420
inputs  ----CCGA-----TGTTGCTT---TCCC---CTGTTGCTAGATTAGCGTCTGCCTCCC----
      3290      3300      3310      3320      3330      3340      3350
      GGAGCCAGCATGGTATGGGAGAGTGCCTGTGAACCTGCCAGGAGCAGGGCCTGGACCAGCAGGCCATGA
      2430      2440      2450      2460      2470
inputs  -----CCTAGTGGAG-----AGGCTGA---TCGC-CAGCT--CTCTGATGCAGGACTCTGGT--
      3360      3370      3380      3390      3400      3410
      ATAGACATACTTGGTGAAGTGAACGGAGACTGAGGATGGCTCTGCTTCCACCGAGG-GAGACACTAGTTG
      2480      2490      2500      2510
inputs  GTTTAGGCTCA--CTCACTATTGGTTTCCTTGGCACAGG-----GTAGTCA----CT-----
      3420      3430      3440      3450      3460      3470      3480
      GCAAAGTGTCTAACCTCCCTTTCCAGCCATTGCTCAAGTCCCCCAGGCTGTGGACATGAGCTGGTGGG
      2520      2530      2540      2550      2560
inputs  CAA---TAAATGTTCC--TCT-----AAAAGCTGAAAAAAAAAAAAAAAAAAGG
      3490      3500      3510      3520      3530      3540      3550
      CAGAATGTTGTTGTTGAAGTCTGATTTTAGATTGATTTTAAAAAAAAAAAAAAAAAAAAAAAAAAGG
      GCGGCCGC
      GCGGCCGC
      3560

```

FIG.35F

	10	20	30	40	50	60	70
inputs	MAPARAGFCPLLLLLLGLWVAEIPVSAKPGMTSSQWFKIQHMQPSQACNSAMKNINKHTKRCKDLNT						

	MV-----LCFPLLLLLLVWGPVCPHAWPKRLTKAHWFEIQHIQPSPLQCNRAMSGINNYAQHCKHQNT						
	10	20	30	40	50	60	

	80	90	100	110	120	130	140
inputs	FLHEPFSSVAATCQTPKIACKNGDKNCHQSHGPVSLTMCKLTSGKYPNCRYKEKRONKSYVVACKPPQKK						

	FLHDSFQNVAAVCDLLSIVCKNRRRHCHQSSKPVNMTDCRLTSGKYPQCRYAAAQYKFFIVACDPPQKS						
	70	80	90	100	110	120	130

	150
inputs	DSQQFHLVPVHLDRL

	DPP-YKLMVPVHLDL
	140 150

FIG.36

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```

inputs  GTCGACCCACGCGTCCGGCTCCCAGCCCACCCCCAACAGACACAGCGTAGCCCGGGCCAGCTCTTAAGG
        AT-----GG

inputs  AGTTCAGGAGTGAGAAGAGGCCCTCAGAGATCTGACAGCCTAGGAGTGGGTGGACACCACCTCAGCCAC
        TG-----CTA---TGCTT---TCCTCTCT---
                10      20

inputs  TGAGCAGGAGTCACAGCACGAAGACCAAGCGCAAAGCGACCCCTGCCCTCCATCCTGACTGCTCCTCCTA
        TTTACTG-----CTGC-----TGGTT-----CTA
                30      40

inputs  AGAGAGATGGCACCAGCCAGAGCAGGATTCTGCCCTTCTGCTGCTTCTGCTGCTGGGCTGTGGGTGG
        TGGG-----GACCAGTG-----TGTCACCTTCA---TGCTT-----GGC-----
                50      60      70

inputs  CAGAGATCCCAGTCAGTGCCAAGCCCAAGGGCATGACCTCATCAGTGGTTTAAATTCAGCACATGCA
        CTAAG---C-GTCT---CA---CCAAGG-C-----TCAC---TGGTTTGAATTCAGCATATACA
                80      90      100      110

inputs  GCCCAGCCCTCAAGCATGCAACTCAGCCATGAAAAACATTAACAAGCACACAAACGGTGCAAAGACCTC
        GCCAAGTCCTCT-----CCA-----ATGCA-----ACAGGGCAATGA-----
        120      130      140      150

inputs  AACACCTTCCTGCAGGAGCCTTTCTCCAGTGTGGCCGCCACCTGCCAGACCCCCAAAATAGCCTGCAAGA
        -----GTGGCATCAAC-----AATTATGCC-----
                160      170

inputs  ATGGCGATAAAAACTGCCACCAGAGCCACGGGCCCGTGTCCCTGACCATGTGTAAAGCTCACCTCAGGGAA
        -----CAG---CAC-----TGTAAAGCA---TCA---A
                180

inputs  GTATCCGAACTGCAGGTACAAAGAGAAGCGACAGAACAAGTCTTACGTAGTGGCCTGTAAGCCTCCCAG
        AATACCTTCTGCATG-AC-----TCTTTC-----CAG
        190      200      210

inputs  AAAAAGGACTCTCAGCAATCCACCTGGTTCTGTACACTTGGACAGAGTCCTTTAGGTTTCCAGACTGG
        AATGTGG---CTGCTGT---CTGT-----GATTTCCT---CAG---
        220      230      240

```

FIG.37A

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```

      710      720      730      740      750      760      770
inputs  CTTGCTCTTTGGCTGACCTTCAATTCCTCTCCAGGACTCCGCACCACTCCCCTACACCCAGAGCATTCT
      .....CATTTGCTG--CAA-AAATC-----GTCG--GCACAAC TGCCA-----CCAGAGC-----
            250            260            270            280

      780      790      800      810      820      830      840
inputs  CTCCCCCTCATCTCTTGGGGCTGTTCTGGTTTCAAGCTCTGCTGGGAGGCTGAAGCTGACACTCTGGTGA
      .....TCAAAG-----CCTG--TCAACAT-GACT---GACTG---CAGACTCACT---
            290            300            310            320

      850      860      870      880      890      900      910
inputs  GCTGAGCTCTAGAGGGATGGCTTTTCATCTTTTGTGCTGTTTCCCAGATGCTTATCCCCAAGAAACA
      .....TCAGGAAAG-----TATCCCAG-----
            330

      920      930      940      950      960      970      980
inputs  GCAAGCTCAGGTCTGTGGGTTCCCTGGTCTATGCCATTGCACATGTCTCCCTGCCCTGGCATTAGGG
      .....TGC-----GCTATAGTG
            340            350

      990      1000      1010      1020      1030      1040      1050
inputs  CAGCATGACAAGGAGAGGAAATAAATGGAAAGGGGGCATATGGGATTTGTGGACACAGCTGTTTCTGTTC
      CTGCT-----GC-----C
            360

      1060      1070      1080      1090      1100      1110      1120
inputs  CTGAAGTAGAAGTCTTCCCCAGCTCTGACGTGGCAGTGAGGTGACCTGAAGGAAAGAAAAATATAAATAA
      CAGTACAAAT--TCTTC-----ATTG
            370

      1130      1140      1150      1160      1170      1180      1190
inputs  ATACCACTTCATATTTGTATAGAATCCTCTAATCCCTTGTGACATAGACTTGACAGGGATTGTATGCCTT
      TTGCCCT-----GTGACC-----CCC-----CT--CAG-----
            380            390

      1200      1210      1220      1230      1240      1250      1260
inputs  CTTTATGGATGAGGAAATTAAGGTTT TAGAAAGCTTAATGAATTAAGAGCTTGTCTAATTAGTTAGTAG
      .....AAGAGC-----
            400

      1270      1280      1290      1300      1310      1320      1330
inputs  CAGAACCTGGACTTGAACCTAGGTCTCCTTGCTCTAAATACAGTGACCTTCTACTCTACCAGTTGCGCA
      ---GACC-----CC-----CC-----CTACAAGTTG---
            410            420

      1340      1350      1360      1370      1380      1390      1400
inputs  AGAAAGAAGTCACTGTTACAGAGGCAAGCGGTGAAGTGAAGTTCCTCATGAAGAAACGAGTGCT
      -----GTTT-CTGT-ACA-----CTTAGATAGTATTCTCT-----
            430            440            450

```

FIG.37B

```

      1410      1420      1430      1440      1450      1460      1470
inputs CTGAAGAGCCAGTTACCCTGTGTTGGCTGCAATAAAGGTCATTACCTCTCTAGCCAAAAAAAAAAAAAA
.....

      1480      1490
inputs AAAAAAAAAAAAAAAAAAAAAAAAAAA
.....AA

```

FIG.37C

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```

      240      250      260      270      280      290      300
AGGATTCTGCCCCCTTCTGCTGCTTCTGCTGCTGGGGCTGTGGGTGGCAGAGATCCCAGTCAGTGCCAAG
GGTGCTATGCTTTCCTTCTTTTACTGCTGCTGTTCTATGGGACCAGTGTGTCCTTCATGCTTGG
      10      20      30      40      50      60      70

      310      320      330      340      350      360      370
CCCAAGGGCATGACCTCATCACAGTGGTTTAAAAATTCAGCACATGCAGCCCAGCCCTCAAGCATGCAACT
CCTAAGCGTCTCACCAGGCTCACTGGTTTGAATTCAGCATATACAGCCAAGTCTCTCCAATGCAACA
      80      90      100      110      120      130      140

      380      390      400      410      420      430      440
CAGCCATGAAAAACATTAACAAGCACAAAAACGGTGCAAAGACCTCAACACCTTCCTGCACGAGCCTTT
GGGCAATGAGTGGCATCAACAATTATGCCAGCACTGTAAGCATCAAAATACCTTTCTGCATGACTCTTT
      150      160      170      180      190      200      210

      450      460      470      480      490      500      510
CTCCAGTGTGGCGCCACCTGCCAGACCCCAAAATAGCCTGCAAGAATGGCGATAAAACTGCCACCA
CCAGAAATGTGGCTGCTGTCTGTGATTGCTCAGCATTTGCTGCAAAAATCGTCGGCACAACCTGCCACCA
      220      230      240      250      260      270      280

      520      530      540      550      560      570      580
GAGCCACGGGCGCGTGTCCCTGACCATGTGTAAGCTCACCTCAGGGAAGTATCCGAACTGCAGGTACAAA
GAGCTCAAAGCCTGTCAACATGACTGACTGCAGACTCACTTCAGGAAAGTATCCCCAGTGGCGCTATAGT
      290      300      310      320      330      340      350

      590      600      610      620      630      640      650
G-AGAAGCGACAGAACAAAGTCTTACGTAGTGGCCTGTAAAGCCTCCCCAGAAAAAGGACTCTCAGCAATTC
GCTGCTGC-CCAGTACAAATTCTTCATTGTTGCCTGTGACCCCTCAGAAAGAGCGACCCCTC-C-TAC
      360      370      380      390      400      410

      660      670      680
CACCTGGTTCCTGTACACTTGGACAGAGTCCTTTAG
AAGTTGGTTCCTGTACACTTAGATAGTATTCTCTAA
      420      430      440      450

```

43.4% identity in 477 aa overlap; score: 746

```

      410      420      430      440      450      460
GGTGCAAAG--ACCTCAACACCTTC-CTGCACGAGCCTTC--TCCAGTGTGGCGCCACCTGCCAGA
GGTGCTATGCTTTCCTTCTTTTACTGCTGCTGTTCTATGGGACCAGTGTGTCCTTCATGCTTGG
      10      20      30      40      50      60      70

```

FIG.38A

470 CC-----CCC AAAATAGCCTGCAAGAATGGCGATAAA-AACTGCCACCAGAGCCACGGGCGCGTGTCC
80 90 100 110 120 130 140
CCTAAGCGTCTCACC AAGGCTCACTGGT TTTGAAATTCAGCATATACAGCC AAGTCTCT- -CCAATGCAA
150 160 170 180 190 200
CTGACCATGTGTAAGCTCACCTCAGGGAAGTATCCGAACTGCAGGTACAAAGAGAAGCGACAGAACAAGT
210 220 230 240 250 260 270
CAGGGCAATGAGTGGCA-TCAACAATTATG- -CCAGCACTGTAAAGCATCAAATACCTTTCTGCATGA
280 290 300 310 320
CTTACGTAGTGGCCTGTAAAGCCTCCCCAGAAAAAGGACT-CTCAGCAAT-TCCACCTGGTTCCTGTACAC
330 340 350 360 370 380 390
CT- -CTTT- -CCAGAATGTGGCTGCTGTCTGTGATTGTCTCAGCAATGTCTGCAAAAATCGTCGGCAC
400 410 420 430 440 450
670 TTGGACAGAGTCCTTTAGGTTTCCAGACTGGCTTGCTCTTTGGCTGACCTTCAATTCCCTCTCCAGGA- -
A- -ACTG- -CCACCAGAGCTCAAAGC- -CTGTCAACATGACTGAC-TGCAGA-CTCACTTCAGGAAA
460 470 480 490 500 510 520 530
---CTCC-GCACCCTCC- -CTACA-CCAGAGCATTCCTTCCCCTCATCTCTTGGGCTGTTC-C
540 550 560 570 580 590 600
GTATCCCAGTGCCTATAGTGTCTGCTGCCAGTACAAATCTTCA- -TTGTTGCTGTGACCCCCCTC
610 620 630 640 650 660 670
330 340 350 360 370 380 390
800 810 820 830 840 850
TG- -GTTCAGCCTCTGCTGGGAGGCTGAAGCTGACACTCTGGTGAGCTGAGCTCTAG
400 410 420 430 440 450
AGAAGAGCGACCCCCCTACAAGTTGGTTCTCTGT-ACACTTAGATAGTATTCTCTAA

TGCAGAGCCTTTCTCCAGTGTGGCCGCCACCTG--CCA-GACCCCCAAAATAGCC--TGCAAAGATGGC
 10 20 30 40 50 60 70
 TGCT-ATGCTTTCCTCTTCTTTTACTGCTGCTGGTCTATGGGGACCAAGTGTGTCCACTTCATGCTTGGC
 80 90 100 110 120 130
 GATAAAACTGCCACCAGAGC-CACGGGCCCGTGTCCCTGACCATGTGTAAGCTCA-CCTCAGGGAAGTA
 140 150 160 170 180
 CTAAGCGTCT--CACCAAGGCTCACTGGTTTGAAATTCAG--CATATACAGCCAAGTCTC-----
 190 200 210 220 230 240 250
 TCCGAA-CTGCAGGTACAAAGAGAAGCGACAGAACAAGTCTTACGTAGTGGCCTGTAAAGCCTCCCAGAA
 260 270 280 290 300
 TCCAATGCAACAGG-GCAATGAGTGGCATC--AACAAATTATGCCAGCA--CTGTAAAGCATC-----A
 310 320 330 340 350
 AAAGGACTCTCAGCAATCCACCTGGTTCCTGTACACTTGGACAGAGTCCCTTTAGGTTTC-CAGACTGGC
 360 370 380 390 400
 AAATACCTTTTCTGCACTACT-CT--TTCCAGAA--TGTTGGCTGCTGTCTGTGATTGCTCAGCAATTGT
 410 420 430 440 450

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laminin_EGF: domain 1 of 4, from 3 to 37: score -1.2, E = 0.59

```

      *->CdCnphGsIsddtCdsddelfgeetGqC1kCkpnvtGrrCdr.CkpG
            + G      d+      ++GqC+ C+ + +G+rC +C +G
mT272    3    ---HASG-----DP-----VHGQCR-CQAGWMGTRCHLPCEG 31
      yyg1psgdpgggC<.*
            ++g      + +C
mT272    32 FWG-----A-NC      37

```

EGF: domain 1 of 4, from 37 to 67: score 19.2, E = 0.1

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyy1syTGkrC<-
            C+ ++ C+ngGtCv+ g      C+C+pG      + G+ C
mT272    37    CSNTCTCKNGGTCVSENG-----NCVCAPG-----FRGPSC 67

```

DSL: domain 1 of 1, from 10 to 67: score -21.2, E = 8.1

```

      *->Wstdkhiggrts1GfnleyrirtCdenYYGsgCnkFCrPrdDafgH
            + ++      + r + C e G+ C++ C      +g+
mT272    10    --HGQCRQCAG---WMGTRCHLPCPEGFWGANCSNTCTCK---NGG 47
      ytCdenGnk1C1eGwkGeyC<.*
            +enGn C++G +G+ C
mT272    48 TCVSENGNCVCAPGFRGPSC      67

```

laminin_EGF: domain 2 of 4, from 41 to 80: score -1.5, E = 0.63

```

      *->CdCnphGsIsddtCdsddelfgeetGqC1kCkpnvtGrrCdr.CkpG
            C+C + G      tC s      e G C+ C p++ G+ C r+C pG
mT272    41    CTCKNGG-----TCVS-----ENGNCV-CAPGFRGPSCQRCPPG 74
      yyg1psgdpgggC<.*
            y      + + C
mT272    75 RY-----GKR--C      80

```

EGF: domain 2 of 4, from 80 to 110: score 11.8, E = 1.9

```

      *->CapnnpCsng.GtCvntpggssdnfggytCeCppGdyy1syTGkrC<-
            C + C+n++ C+++ g      Tc C G      +tG++C
mT272    80    CVQC-KCNNNhSSCHPSDG-----TCSCLAG-----WTGPDC 110

```

FIG.39A

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laminin_EGF: domain 3 of 4, from 83 to 123: score 25.6, E = 0.0012

```

          *->CdCnphGalsddtCdsddelfgeetGqClkCkpnvtGrrC.drCkpG
          C Cn++ ++C++ +G C+ C+ + tG++C++ C pG
mT272    83  CKCNNH-...SSCHP-.....SDGTCS-CLAGWTGPDCsEACPPG 117

          yyglpsgdpqgqC<-*
          ++gl      C
mT272    118 HWGL-.....KC      123

```

EGF: domain 3 of 4, from 123 to 153: score 27.3, E = 0.00036

```

          *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyy1syTGkrC<-
          C++++ C++gGtc++ g +C+C+pG +tG++c
mT272    123  CSQLCQCHHGGTCHPQDG-.....SCICTPG-....WTGPNC 153

```

laminin_EGF: domain 4 of 4, from 127 to 172: score -5.5, E = 1.4

```

          *->CdCnphGs1sddtCdsddelfgeetGqClkCkpnvtGrrC.drCkpG
          C+C++ G      tC++      G C C p+ tG++C + C p
mT272    127  CQCHHGG-....TCHP-.....QDGSCI-CTPGWTGPNC1EGCPPR 160

          yyglpsg.dpgqgC<-*
          +g ++++ + +C
mT272    161 MFG-VNCsQLC-QC      172

```

EGF: domain 4 of 4, from 166 to 196: score 4.5. E = 5.8

```

          *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyy1syTGkrC<-
          C++++ C+ g C++ g      C+CppG +G +C
mT272    166  CSQLCQCDLGEMCHPETG-.....ACVCPPG-....HSGADC 196

```

FIG.39B

PFAM

EGF-like REPEATS AND FN-3 like REPEATS

9945



Cys
Ngly

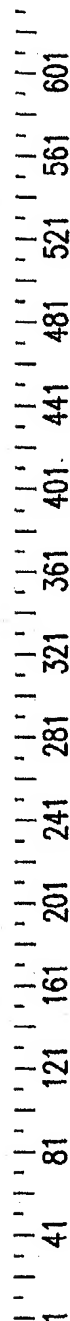


FIG.40

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```

      *->CaPnnpCsnqGtCvntpggssdntgggtCeCppGayylsyGkrC<-
      C+++ C+ngG C g +C+C+pG y+G+rC
ratT272 18 IECRCHNGGLCDRFTG-----QCHCAPG-----YIGDPRC 48

```

laminin_EGF: domain 1 of 11, from 22 to 61: score 12.3, E = 0.038

```

      *->CdCnphGsIsddtCdsddelfgeetGqClkCkpnvtGrrC.drCkpG
      C C++ G Cd+ +tGqC+ C p++ G+rC+++C G
ratT272 22 CRCHNGG-----LCDR-----FTGQCH-CAPGYIGDRCrEECPVG 55

      yyglpsgdpgggC<-*
      +g g+C
ratT272 56 RFG-----QDC 61

```

EGF: domain 2 of 11, from 61 to 91: score 18.3, E = 0.18

```

      *->CapnnpCsnqGtCvntpggssdnfggytCeCppGdyylsyGkrC<-
      Ca+++ C q++C + g C C +G +tG+rC
ratT272 61 CAETCDCAPGARCFPANG-----ACLCEHG-----FTGDRC 91

```

laminin_EGF: domain 2 of 11, from 65 to 105: score 4.0, E = 0.2

```

      *->CdCnphGsIsddtCdsddelfgeetGqClkCkpnvtGrrCdr..Ckp
      CdC p + +C + G+C1 C +++tG+rC ++ C +
ratT272 65 CDCAPGA-----RCFP-----ANGACL-CEHGFTGDRCter1CPD 98

      GyyglpsgdpgggC<-*
      G yg1 +C
ratT272 99 GRYGL-----SC 105

```

EGF: domain 3 of 11, from 105 to 137: score 4.1, E = 9.6

```

      *->CapnnpCsnq..GtCvntpggssdnfggytCeCppGdyylsyGkrC
      C++++ C+ ++ C++ +g +C C+pG ++G +C
ratT272 105 CQDPCTCDPEhsLSCHPMHG-----ECSCQPG-----WAGLHC 137

```

laminin_EGF: domain 3 of 11, from 109 to 150: score 13.1, E = 0.032

```

      *->CdCnphGsIsddtCdsddelfgeetGqClkCkpnvtGrrCdr.CkpG
      C+C+p sls C++ ++G+C+ C+p ++G +C+++C
ratT272 109 CTCDPEHSLS---CHP-----MHGECS-CQPGWAGLHCNEscP-- 142

      yyglpsgdpgggC<-*
      ++ +g gC
ratT272 143 --QD---THGAGC 150

```

FIG.41A

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EGF: domain 4 of 11, from 150 to 180: score 27.7, E = 0.00026

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
      C++++ C++gG+C+  g          C+C+pG      ytG++C
ratT272  150  CQEHCLCLHGGVCLADSG-----LCRCAPG-----YTGPNC  180

```

laminin_EGF: domain 4 of 11, from 154 to 193: score 8.4, E = 0.084

```

      *->CdCnphGsIsddtCdsdelfgeetGqClkCkpnvtGrrC.drCkpG
      C C +hg      + C          +G C+ C p++tG++C + C p+
ratT272  154  CLC-LHG----GVCLA-----DSGLCR-CAPGYTGPNaNLCPPN  187

      yyglpsgdpgqgC<-*
      *yg          +C
ratT272  188 TYGI-----NC      193

```

EGF: domain 5 of 11, from 193 to 223: score 10.6, E = 2.5

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
      C++++ C n      C ++ g          tC+C++G      ++ +C
ratT272  193  CSSHCSCENAIAACSPVDG-----TCICKEG-----WQRGNC  223

```

laminin_EGF: domain 5 of 11, from 197 to 236: score 0.7, E = 0.4

```

      *->CdCnphGsIsddtCdsdelfgeetGqClkCkpnvtGrrCdr.CkpG
      C C ++      C +          + G C CK++ + +C +C pG
ratT272  197  CSCENAI-----ACSP-----VDGTCTI-CKEGWQRGNCSVpCPPG  230

      yyglpsgdpgqgC<-*
      ++g+          +C
ratT272  231 TWG-----SC      236

```

EGF: domain 6 of 11, from 236 to 266: score 11.8, E = 1.9

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
      C+  + C + G+C +  g          C+C+pG      + G +C
ratT272  236  CNASCQCAHEGVCSPTG-----ACTCTPG-----WRGVHC  266

```

laminin_EGF: domain 6 of 11, from 240 to 279: score -2.2, E = 0.73

```

      *->CdCnphGsIsddtCdsdelfgeetGqClkCkpnvtGrrCdr.CkpG
      C+C + G      C +          tG+C C p+ G +C +C G
ratT272  240  CQCAHEG-----VCSP-----QTGACT-CTPGWRGVHCQLpCPKG  273

      yyglpsgdpgqgC<-*
      +g          +gC
ratT272  274 QFG-----EGC      279

```

FIG.41B

DSL: domain 1 of 1, from 246 to 309: score -19.4, E = 5.2

```

      *->WstdkhiggrtslGfnleyrivrvtCdenYYGegCnkFCrPrdDafgH
            + ++++g+ t      +++ C + +GegC+ C+      H
ratT272  246  GVCSPQTGACTCTPGQRGVHCQLPCPKGQFGEGCASVCDCH 287

            yt.Cd.enGnk1c1eGwkGeyC<-*
            + +Cd+ +G +C +GW+G C
ratT272  288 SDgCDpVHGHCRCQAGWMGTRC      309

```

EGF: domain 7 of 11, from 279 to 309: score 7.0, E = 5.3

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
            Ca+ + C++ C +++g      +C+C+ G      + G rC
ratT272  279  CASVCDCHSDGCDPVHG-----HCRCQAG-----WMGTRC 309

```

laminin_EGF: domain 7 of 11, from 283 to 322: score 12.7, E = 0.035

```

      *->CdCnphGs1sddtCdsddelfgeetGqC1kCkpnvtGrrCdr.CkpG
            CdC+ h+ d Cd+      ++G+C+ C+ + +G+rC +C +G
ratT272  283  CDCD-HS----DGCDP-----VHGHCRCQAGQMGRCHLPCEG 316

            yyg1psgdpqgqC<-*
            ++g      + +C
ratT272  317 FWG-----A-NC      322

```

EGF: domain 8 of 11, from 322 to 352: score 17.3, E = 0.38

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
            C+ + C+ngGtCv+ g      C+C+pG      + G+ C
ratT272  322  CSNACTCKNGGTCTVPENG-----NCVCAPG-----FRGPSC 352

```

laminin_EGF: domain 8 of 11, from 326 to 365: score -1.8, E = 0.67

```

      *->CdCnphGs1sddtCdsddelfgeetGqC1kCkpnvtGrrCdr.CkpG
            C+C + G      tC +      e G C+ C p++ G+ C r+C pG
ratT272  326  CTCKNGG-----TCVP-----ENGNCV-CAPGFRGPSCQRpCPPG 359

            yyg1psgdpqgqC<-*
            y      + + C
ratT272  360 RY-----GKR-C      365

```

EGF: domain 9 of 11, from 365 to 394: score 18.3, E = 0.18

```

      *->CapnnpCsngGtCvntpggssdnfggytCeCppGdyylsyTGkrC<-
            C p C+n+ C+++ g      tC C G      +tG++C
ratT272  365  CVPC-KCNNHSSCHPSDG-----TCSCLAG-----WTGPDC 394

```

FIG.41C

laminin_EGF: domain 9 of 11, from 368 to 407: score 24.0, E = 0.0034

```

      *->CdCnphGs1sddtCdsddefgeetGqC1kCkpnvtGrrC.drCkpG
      C Cn+h+ +C++ + G C+ + + tG++C++ C pG
ratT272 368 CKCNNHS-----SCHP-----SDGTCS-CLAGWTGPDCsESCPPG 401
      yyglpsgdpgqgC<-*
      ++gl C
ratT272 402 HWGL-----KC 407

```

EGF: domain 10 of 11, from 407 to 437: score 24.0, E = 0.035

```

      *->CapnnpCsngGtCvntpggssdnfgytCeCppGdyylsyTGkrC<-
      C++++ C++g+tC++ g +C+C pG +tG++C
ratT272 407 CSQPCQCHHGATCHPQDG-----SCVCIPG-----WTGPNC 437

```

laminin_EGF: domain 10 of 11, from 407 to 437: score 6.5, E = 0.12

```

      *->CdCnphGs1sddtCdsddefgeetGqC1kCkpnvtGrrCdrCkpGy
      C+C++ + tC++ G C+ C p+ tG++C +
ratT272 411 CQCHGA-----TCHP-----QDGSCV-CIPGWTGPNCSE----- 439
      yg1psgdpgqgC<-*
      g ps+++g++C
ratT272 440 -GCPSRMFGVNC 450

```

EGF: domain 11 of 11, from 450 to 480: score 8.7, E = 3.7

```

      *->CapnnpCsngGtCvntpggssdnfgytCeCppGdyylsyTGkrC<-
      C++++ C+ g C++ g C+CppG +G +C
ratT272 450 CSQLCQCDPGEMCHPETG-----ACVCPPG-----HSGAHC 480

```

laminin_EGF: domain 11 of 11, from 454 to 489: score -6.3, E = 1.7

```

      *->CdCnphGs1sddtCdsddefgeetGqC1kCkpnvtGrrCdrCkpGy
      C+C+p G + C++ etG+C+ C p+ +G +C
ratT272 454 CQCDP-G-----EMCHP-----ETGACV-CPPGHSGAHC-----K 481
      yg1psgdpgqgC<-*
      g + ++
ratT272 482 VGSQE-SFT--- 489

```

FIG.41D

SEQUENCE LISTING

<110> Millennium Pharmaceuticals, Inc.

<120> MEMBRANE-ASSOCIATED AND SECRETED PROTEINS AND USES THEREOF

<130> 7853-206-228

<150> 09/345,464

<151> 1999-06-30

<160> 148

<170> FastSEQ for Windows Version 3.0

<210> 1

<211> 3284

<212> DNA

<213> Homo sapiens

<220>

<221> CDS

<222> (1222) ... (1944)

<400> 1

```

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gattttgttt tctttcatcc cttttcccaa gcagtttatt atgaaaattt tcaaacatac      120
agcaatgttg agaaaatttt acagtaaatg cctataccca ttacctaaat tttaccatta      180
acattttacc ctgctggcat tattgtgctt atccatctac gtatccctct ctcccttcat      240
tggtgtatatt ctaagtaaat tgtaggcctc agtacacttc cttctgaatt cttcagcatg      300
cacaacagta ttatattcca tttttaaaag agcaattctt gatagattta tatagttttg      360
taaaatgttc atatagagct acaaatttta tctttttggt tcttattgta tgtctagggg      420
cctgaagggg atgctggcat tgttgggata tcaggtccta aaggctctat tggacacaga      480
ggaaacactg gtccccttgg cagagaaggt ataataggcc caacaggtag aactggaccc      540
agagggtgaaa agggcttttag aggtgaaact ggtcctcaag gaccaagagg tcaaccaggg      600
cctccaggtc cactgggagc accaggccca agaaagcaaa tggatatcaa tgctgctatt      660
caagccttga ttgaatcaaa tactgcccta cagatggagg taacatatct ggttttattt      720
atattggcac tgtctctcaa tataccaatt aaacagagaa aatttttga ggccaaaatg      780
tgacattatc tcaaagattg tattttaaac agattgaaaa tgtgaaacca ttctcaagaa      840
caaagtaagt gattttggta taattaaaca gaaatatatg cgtaggatgt tttgtaagga      900
aaacatttaa atcaaaaatt tagtactgtt atttgaagg aatttggtag tatccaagaa      960
agtagttaaa tgaggtttag catgtttctt aaaatgagat atatataatta tcaactactca    1020
tttattttaa ctctaattgat tcaatgtgta atttaaaaaa cataatacag tagacatagc    1080
aattccttatg ttagcttgaa aactaaactt gcaaatgtga atttaacctc tttaaaagat    1140
taagggttatt aaagcataca catatgccta tgcttaaata taaactgttc tttacattct    1200
actcacaact tactacacat a atg gaa aca cat tct tct cct gcc ttg gcc    1251
                               Met Glu Thr His Ser Ser Pro Ala Leu Ala
                               1             5             10

cat gtt ggt cct cag gat ttt ttt gtt tat ata att ctt atg atg act      1299
His Val Gly Pro Gln Asp Phe Phe Val Tyr Ile Ile Leu Met Met Thr
                               15             20             25

tgg cag agc tac cag aat act gaa gtg act tta att gac cac agt gaa      1347
Trp Gln Ser Tyr Gln Asn Thr Glu Val Thr Leu Ile Asp His Ser Glu
                               30             35             40

```

gag ata ttc aaa acc ctg aac tac ctt agc aat tta ttg cac agc atc	1395
Glu Ile Phe Lys Thr Leu Asn Tyr Leu Ser Asn Leu Leu His Ser Ile	
45 50 55	
aag aat cct ctt ggc aca cga gat aac cca gca cga atc tgc aaa gat	1443
Lys Asn Pro Leu Gly Thr Arg Asp Asn Pro Ala Arg Ile Cys Lys Asp	
60 65 70	
tta ctt aac tgt gaa caa aaa gta tca gat gga aaa tac tgg att gac	1491
Leu Leu Asn Cys Glu Gln Lys Val Ser Asp Gly Lys Tyr Trp Ile Asp	
75 80 85 90	
cca aat ctt ggc tgt cct tca gat gcc att gag gtt ttc tgc aat ttc	1539
Pro Asn Leu Gly Cys Pro Ser Asp Ala Ile Glu Val Phe Cys Asn Phe	
95 100 105	
agt gct ggt ggc cag aca tgc tta cct cct gtt tct gta aca aag ttg	1587
Ser Ala Gly Gly Gln Thr Cys Leu Pro Pro Val Ser Val Thr Lys Leu	
110 115 120	
gag ttt gga gtt ggg aaa gtc cag atg aac ttc ctt cat tta ctg agt	1635
Glu Phe Gly Val Gly Lys Val Gln Met Asn Phe Leu His Leu Leu Ser	
125 130 135	
tcg gaa gcc acc cat atc atc acc att cac tgt cta aac acc cca agg	1683
Ser Glu Ala Thr His Ile Ile Thr Ile His Cys Leu Asn Thr Pro Arg	
140 145 150	
tgg aca agc aca caa aca agt ggc cca gga ttg cct att ggt ttc aag	1731
Trp Thr Ser Thr Gln Thr Ser Gly Pro Gly Leu Pro Ile Gly Phe Lys	
155 160 165 170	
gga tgg aat ggc cag att ttt aaa gta aac act cta ctt gaa cct aaa	1779
Gly Trp Asn Gly Gln Ile Phe Lys Val Asn Thr Leu Leu Glu Pro Lys	
175 180 185	
gtg ctt tca gat gac tgc aag att caa gat ggc agc tgg cat aag gca	1827
Val Leu Ser Asp Asp Cys Lys Ile Gln Asp Gly Ser Trp His Lys Ala	
190 195 200	
aca ttt ctt ttt cac acc cag gaa cct aat caa ctt cca gtg att gaa	1875
Thr Phe Leu Phe His Thr Gln Glu Pro Asn Gln Leu Pro Val Ile Glu	
205 210 215	
gta caa aaa ctt cct cat ctc aaa act gaa cga aag tat tac att gac	1923
Val Gln Lys Leu Pro His Leu Lys Thr Glu Arg Lys Tyr Tyr Ile Asp	
220 225 230	
agc agt tct gta tgc ttt ctg taaagtctct gaattagttc cgaattcagg	1974
Ser Ser Ser Val Cys Phe Leu	
235 240	
ctgttggcca ggtaattgct gcagagggag aaataagaca gacagataca gtcattatga	2034
aatgcattgta ataaagcatt ggctaaatct taaagaatct caggaagaac agacttcctc	2094
ctaagaagga gaaaaggcat ttttaaagga ctatgattga taaagtattt aattctttta	2154
aaaattatat tcatctcagc tttcttagag aattccctag aactaaaaat ttataaatat	2214
ggaattcttc aggttatctt atatttttga ctgagtgcgt agtaccatt agacagctgg	2274
agatgcagag cactatggag caatactggc taatgcttcc agatgtgcac tgcttctgtc	2334

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taaaaattac aagccacagt ctaatatgtc ttattttcca aaacaotaag ctgtattcag 2394
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tgtgttgctt ggtgctcttt cgaaaacaag gtgcttatgg ctttcataga ctatttctct 2514
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gttagtactt attttaattt gtttggtcac acacttaata acacatgaaa ctatttatgt 2634
gaagtccttg ttttatttta aaattctctt tgtgtatttg gaatcaaagc cagcacattg 2694
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gttgtgcoat actgttttta aagttcatga tcatctggaa tgatacttag tgtatatata 2874
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ctttatatat ctgctttgta gaaattatat gttttgtagt attcattgat tttctttcac 2994
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tccagaaaaa aaaaagtctt ttcccattta aaataggctc agccagttca atgtcgctt 3114
gttatcagag aaatattagt tcaatactga aagaaaaata ttataacctt tggtatctag 3174
aaaagcttgt tcatccatta taaatatatc tttagccaca gcaaaccaca cttaacctat 3234
ctataataaa aatgtgcttt aaataaaaaa aaaaaaaaaa agggcgccgc 3284

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<210> 2
<211> 241
<212> PRT
<213> Homo sapiens

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1 5 10 15
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20 25 30
Thr Glu Val Thr Leu Ile Asp His Ser Glu Glu Ile Phe Lys Thr Leu
35 40 45
Asn Tyr Leu Ser Asn Leu Leu His Ser Ile Lys Asn Pro Leu Gly Thr
50 55 60
Arg Asp Asn Pro Ala Arg Ile Cys Lys Asp Leu Leu Asn Cys Glu Gln
65 70 75 80
Lys Val Ser Asp Gly Lys Tyr Trp Ile Asp Pro Asn Leu Gly Cys Pro
85 90 95
Ser Asp Ala Ile Glu Val Phe Cys Asn Phe Ser Ala Gly Gly Gln Thr
100 105 110
Cys Leu Pro Pro Val Ser Val Thr Lys Leu Glu Phe Gly Val Gly Lys
115 120 125
Val Gln Met Asn Phe Leu His Leu Leu Ser Ser Glu Ala Thr His Ile
130 135 140
Ile Thr Ile His Cys Leu Asn Thr Pro Arg Trp Thr Ser Thr Gln Thr
145 150 155 160
Ser Gly Pro Gly Leu Pro Ile Gly Phe Lys Gly Trp Asn Gly Gln Ile
165 170 175
Phe Lys Val Asn Thr Leu Leu Glu Pro Lys Val Leu Ser Asp Asp Cys
180 185 190
Lys Ile Gln Asp Gly Ser Trp His Lys Ala Thr Phe Leu Phe His Thr
195 200 205
Gln Glu Pro Asn Gln Leu Pro Val Ile Glu Val Gln Lys Leu Pro His
210 215 220
Leu Lys Thr Glu Arg Lys Tyr Tyr Ile Asp Ser Ser Ser Val Cys Phe
225 230 235 240
Leu

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<210> 3
<211> 723

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<212> DNA
<213> Homo sapiens

<400> 3

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ataattctta tgatgacttg gcagagctac cagaatactg aagtgaacttt aattgaccac	120
agtgaagaga tattcaaaac cctgaactac cttagcaatt tattgcacag catcaagaat	180
cctcttggca cactgagataa cccagcacga atctgcaaag atttacttaa ctgtgaacaa	240
aaagtatcag atggaaaata ctggattgac ccaaattctt gctgtccttc agatgccatt	300
gaggttttct gcaatttcag tgcctgggtggc cagacatgct tacctcctgt ttctgtaaca	360
aagttggagt ttggagttgg gaaagtccag atgaacttcc ttcatttact gagttcggaa	420
gccaccata tcatcaccat tcaactgtcta aacaccccaa ggtggacaag cacacaaaca	480
agtggcccag gattgocctat tggtttcaag ggatggaatg gccagatttt taaagtaaac	540
actctacttg aacctaaaagt gcttcoagat gactgcaaga ttcaagatgg cagctggcat	600
aaggcaacat ttctttttca ccccaggaa cctaatacaac ttccagtgat tgaagtacaa	660
aaacttcctc atctcaaaac tgaacgaaag tattacattg acagcagttc tgtatgcttt	720
ctg	723

<210> 4
<211> 3169
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (57)...(1568)

<400> 4

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acg ccg agc ccc ctg ttg ctg ctc ctg ctg ccg ccg ctg ctg ctg ggg	107
Thr Pro Ser Pro Leu Leu Leu Leu Leu Leu Pro Pro Leu Leu Leu Gly	
5 10 15	
gcc ttc ccg ccg gcc gcc gcc gcc cga ggc ccc cca aag atg gcg gac	155
Ala Phe Pro Pro Ala Ala Ala Ala Arg Gly Pro Pro Lys Met Ala Asp	
20 25 30	
aag gtg gtc cca ccg cag gtg gcc ccg ctg ggc cgc act gtg ccg ctg	203
Lys Val Val Pro Arg Gln Val Ala Arg Leu Gly Arg Thr Val Arg Leu	
35 40 45	
cag tgc cca gtg gag ggg gac ccg ccg ccg ctg acc atg tgg acc aag	251
Gln Cys Pro Val Glu Gly Asp Pro Pro Pro Leu Thr Met Trp Thr Lys	
50 55 60 65	
gat ggc cgc acc atc cac agc ggc tgg agc cgc ttc cgc gtg ctg ccg	299
Asp Gly Arg Thr Ile His Ser Gly Trp Ser Arg Phe Arg Val Leu Pro	
70 75 80	
cag ggg ctg aag gtg aag cag gtg gag ccg gag gat gcc gcc gtg tac	347
Gln Gly Leu Lys Val Lys Gln Val Glu Arg Glu Asp Ala Gly Val Tyr	
85 90 95	
gtg tgc aag gcc acc aac gcc ttc gcc agc ctg agc gtc aac tac acc	395
Val Cys Lys Ala Thr Asn Gly Phe Gly Ser Leu Ser Val Asn Tyr Thr	

100	105	110	
ctc gtc gtg ctg gat gac att agc cca ggg aag gag agc ctg ggg ccc Leu Val Val Leu Asp Asp Ile Ser Pro Gly Lys Glu Ser Leu Gly Pro 115 120 125			443
gac agc tcc tct ggg ggt caa gag gac ccc gcc agc cag cag tgg gca Asp Ser Ser Ser Gly Gly Gln Glu Asp Pro Ala Ser Gln Gln Trp Ala 130 135 140 145			491
cga ccg cgc ttc aca cag ccc tcc aag atg agg cgc cgg gtg atc gca Arg Pro Arg Phe Thr Gln Pro Ser Lys Met Arg Arg Arg Val Ile Ala 150 155 160			539
cgg ccc gtg ggt agc tcc gtg cgg ctc aag tgc gtg gcc agc ggg cac Arg Pro Val Gly Ser Ser Val Arg Leu Lys Cys Val Ala Ser Gly His 165 170 175			587
cct cgg ccc gac atc acg tgg atg aag gac gac cag gcc ttg acg cgc Pro Arg Pro Asp Ile Thr Trp Met Lys Asp Asp Gln Ala Leu Thr Arg 180 185 190			635
cca gag gcc gct gag ccc agg aag aag aag tgg aca ctg agc ctg aag Pro Glu Ala Ala Glu Pro Arg Lys Lys Lys Trp Thr Leu Ser Leu Lys 195 200 205			683
aac ctg cgg ccg gag gac agc ggc aaa tac acc tgc cgc gtg tcg aac Asn Leu Arg Pro Glu Asp Ser Gly Lys Tyr Thr Cys Arg Val Ser Asn 210 215 220 225			731
cgc gcg ggc gcc atc aac gcc acc tac aag gtg gat gtg atc cag cgg Arg Ala Gly Ala Ile Asn Ala Thr Tyr Lys Val Asp Val Ile Gln Arg 230 235 240			779
acc cgt tcc aag ccc gtg ctc aca ggc acg cac ccc gtg aac acg acg Thr Arg Ser Lys Pro Val Leu Thr Gly Thr His Pro Val Asn Thr Thr 245 250 255			827
gtg gac ttc ggg ggg acc acg tcc ttc cag tgc aag gtg cgc agc gac Val Asp Phe Gly Gly Thr Thr Ser Phe Gln Cys Lys Val Arg Ser Asp 260 265 270			875
gtg aag ccg gtg atc cag tgg ctg aag cgc gtg gag tac ggc gcc gag Val Lys Pro Val Ile Gln Trp Leu Lys Arg Val Glu Tyr Gly Ala Glu 275 280 285			923
ggc cgc cac aac tcc acc atc gat gtg ggc ggc cag aag ttt gtg gtg Gly Arg His Asn Ser Thr Ile Asp Val Gly Gly Gln Lys Phe Val Val 290 295 300 305			971
ctg ccc acg ggt gac gtg tgg tcg cgg ccc gac ggc tcc tac ctc aat Leu Pro Thr Gly Asp Val Trp Ser Arg Pro Asp Gly Ser Tyr Leu Asn 310 315 320			1019
aag ctg ctc atc acc cgt gcc cgc cag gac gat gcg ggc atg tac atc Lys Leu Leu Ile Thr Arg Ala Arg Gln Asp Asp Ala Gly Met Tyr Ile 325 330 335			1067

tgc ctt ggc gcc aac acc atg ggc tac agc ttc cgc agc gcc ttc ctc	1115
Cys Leu Gly Ala Asn Thr Met Gly Tyr Ser Phe Arg Ser Ala Phe Leu	
340 345 350	
acc gtg ctg cca gac cca aaa ccg cca ggg cca cct gtg gcc tcc tgc	1163
Thr Val Leu Pro Asp Pro Lys Pro Pro Gly Pro Pro Val Ala Ser Ser	
355 360 365	
tcc tcg gcc act agc ctg ccg tgg ccc gtg gtc atc ggc atc cca gcc	1211
Ser Ser Ala Thr Ser Leu Pro Trp Pro Val Val Ile Gly Ile Pro Ala	
370 375 380 385	
ggc gct gtc ttc atc ctg ggc acc ctg ctc ctg tgg ctt tgc cag gcc	1259
Gly Ala Val Phe Ile Leu Gly Thr Leu Leu Leu Trp Leu Cys Gln Ala	
390 395 400	
cag aag aag ccg tgc acc ccc gcg cct gcc cct ccc ctg cct ggg cac	1307
Gln Lys Lys Pro Cys Thr Pro Ala Pro Ala Pro Pro Leu Pro Gly His	
405 410 415	
cgc ccg ccg ggg acg gcc cgc gac cgc agc gga gac aag gac ctt ccc	1355
Arg Pro Pro Gly Thr Ala Arg Asp Arg Ser Gly Asp Lys Asp Leu Pro	
420 425 430	
tcg ttg gcc gcc ctc agc gct ggc cct ggt gtg ggg ctg tgt gag gag	1403
Ser Leu Ala Ala Leu Ser Ala Gly Pro Gly Val Gly Leu Cys Glu Glu	
435 440 445	
cat ggg tct ccg gca gcc ccc cag cac tta ctg ggc cca ggc cca gtt	1451
His Gly Ser Pro Ala Ala Pro Gln His Leu Leu Gly Pro Gly Pro Val	
450 455 460 465	
gct ggc cct aag ttg tac ccc aaa ctc tac aca gac atc cac aca cac	1499
Ala Gly Pro Lys Leu Tyr Pro Lys Leu Tyr Thr Asp Ile His Thr His	
470 475 480	
aca cac aca cac tct cac aca cac tca cac gtg gag ggc aag gtc cac	1547
Thr His Thr His Ser His Thr His Ser His Val Glu Gly Lys Val His	
485 490 495	
cag cac atc cac tat cag tgc tagacggcac cgtatctgca gtgggcacgg	1598
Gln His Ile His Tyr Gln Cys	
500	
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ctggacacac acacacagac acacacactg cctggatgca tgtatgcaca cacatgcgcg	1778
cacacgtgct cctgaaggc acacgtacgc acacacgcac atgcacagat atgccgctg	1838
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gatattgcct ggacacacac acacacacgt gtgcacagat atgctgtctg gacacgcaca	2138
cacatgcaga tatgtgtgct ggacacacac ttccagacac acgtgcacag gcgcagatat	2198
gctgcctgga cacacgcaga tatgtgtctt agtcacacac acacgcagac atgtgttccg	2258
gacacacaca cgcattgcaca gatattgctgt ccggacacac acacgcacgc agatatgctg	2318
cctggacaca cacacagata atgtgtgctc aacactcaca cacgtgcaga tattgctgtg	2378
acacacacat gtgcacagat atgtgtgtctg gacatgcaca cacgtgcaga tatgtgtctc	2438

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ggatacacac gcaacgcacac atgcagatat gctgcctggg cacacacttc cggacacaca 2498
tgacacacaca ggtgcagata tgctgcctgg acacacgcag actgacgtgc ttttgggagg 2558
gtgtgccgtg aagcctgcag tacgtgtgcc gtgaggctca tagttgatga gggactttcc 2618
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agggcgggccg c 3169

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<210> 5
<211> 504
<212> PRT
<213> Homo sapiens

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<400> 5
Met Thr Pro Ser Pro Leu Leu Leu Leu Leu Leu Pro Pro Leu Leu Leu
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Gly Ala Phe Pro Pro Ala Ala Ala Ala Arg Gly Pro Pro Lys Met Ala
20 25 30
Asp Lys Val Val Pro Arg Gln Val Ala Arg Leu Gly Arg Thr Val Arg
35 40 45
Leu Gln Cys Pro Val Glu Gly Asp Pro Pro Pro Leu Thr Met Trp Thr
50 55 60
Lys Asp Gly Arg Thr Ile His Ser Gly Trp Ser Arg Phe Arg Val Leu
65 70 75 80
Pro Gln Gly Leu Lys Val Lys Gln Val Glu Arg Glu Asp Ala Gly Val
85 90 95
Tyr Val Cys Lys Ala Thr Asn Gly Phe Gly Ser Leu Ser Val Asn Tyr
100 105 110
Thr Leu Val Val Leu Asp Asp Ile Ser Pro Gly Lys Glu Ser Leu Gly
115 120 125
Pro Asp Ser Ser Ser Gly Gly Gln Glu Asp Pro Ala Ser Gln Gln Trp
130 135 140
Ala Arg Pro Arg Phe Thr Gln Pro Ser Lys Met Arg Arg Arg Val Ile
145 150 155 160
Ala Arg Pro Val Gly Ser Ser Val Arg Leu Lys Cys Val Ala Ser Gly
165 170 175
His Pro Arg Pro Asp Ile Thr Trp Met Lys Asp Asp Gln Ala Leu Thr
180 185 190
Arg Pro Glu Ala Ala Glu Pro Arg Lys Lys Lys Trp Thr Leu Ser Leu
195 200 205
Lys Asn Leu Arg Pro Glu Asp Ser Gly Lys Tyr Thr Cys Arg Val Ser
210 215 220
Asn Arg Ala Gly Ala Ile Asn Ala Thr Tyr Lys Val Asp Val Ile Gln
225 230 235 240
Arg Thr Arg Ser Lys Pro Val Leu Thr Gly Thr His Pro Val Asn Thr
245 250 255
Thr Val Asp Phe Gly Gly Thr Thr Ser Phe Gln Cys Lys Val Arg Ser
260 265 270
Asp Val Lys Pro Val Ile Gln Trp Leu Lys Arg Val Glu Tyr Gly Ala
275 280 285
Glu Gly Arg His Asn Ser Thr Ile Asp Val Gly Gly Gln Lys Phe Val
290 295 300

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<210> 6
<211> 1512
<212> DNA
<213> Homo sapiens
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8

cactatcagt gc

1512

<210> 7
<211> 1074
<212> DNA
<213> Mus musculus

<220>
<221> CDS
<222> (3)...(626)

<221> modified_base
<222> all "n" positions
<223> n=a, c, g, or t

<400> 7
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Arg Val Arg Pro Thr Gly Asp Val Trp Ser Arg Pro Asp Gly Ser
1 5 10 15

tac ctc aac aag ctg ctc atc tct cgg gcc cgc cag gat gat gct ggc 95
Tyr Leu Asn Lys Leu Ile Ser Arg Ala Arg Gln Asp Asp Ala Gly
20 25 30

atg tac atc tgc cta ggt gca aat acc atg ggc tac agt ttc cgt agc 143
Met Tyr Ile Cys Leu Gly Ala Asn Thr Met Gly Tyr Ser Phe Arg Ser
35 40 45

gcc ttc ctc act gta tta cca gac ccc aaa cct cca ggg cct cct atg 191
Ala Phe Leu Thr Val Leu Pro Asp Pro Lys Pro Pro Gly Pro Pro Met
50 55 60

gct tct tca tcg tca tcc aca agc ctg cca tgg cct gtg gtg atc ggc 239
Ala Ser Ser Ser Ser Thr Ser Leu Pro Trp Pro Val Val Ile Gly
65 70 75

atc cca gct ggt gct gtc ttc atc cta ggc act gtg ctg ctc tgg ctt 287
Ile Pro Ala Gly Ala Val Phe Ile Leu Gly Thr Val Leu Leu Trp Leu
80 85 90 95

tgc cag acc aag aag aag cca tgt gcc cca gca tct aca ctt cct gtg 335
Cys Gln Thr Lys Lys Lys Pro Cys Ala Pro Ala Ser Thr Leu Pro Val
100 105 110

cct ggg cat cgt ccc cca ggg aca tcc cga gaa cgc agt ggt gac aag 383
Pro Gly His Arg Pro Pro Gly Thr Ser Arg Glu Arg Ser Gly Asp Lys
115 120 125

gac ctg ccc tca ttg gct gtg ggc ata tgt gag gag cat gga tcc gcc 431
Asp Leu Pro Ser Leu Ala Val Gly Ile Cys Glu Glu His Gly Ser Ala
130 135 140

atg gcc ccc cag cac atc ctg gcc tct ggc tca act gct ggc ccc aag 479
Met Ala Pro Gln His Ile Leu Ala Ser Gly Ser Thr Ala Gly Pro Lys
145 150 155

ctg tac ccc aag cta tac aca gat gtg cac aca cac aca cat aca cac 527
Leu Tyr Pro Lys Leu Tyr Thr Asp Val His Thr His Thr His Thr His

160 165 170 175

acc tgc act cac acg ctc tca tgt tgg agg gca agg ttc atc aac acc 575
 Thr Cys Thr His Thr Leu Ser Cys Trp Arg Ala Arg Phe Ile Asn Thr
 180 185 190

agc atg tcc act atc agt gct aaa tac agc gaa tct cca agc act gtg 623
 Ser Met Ser Thr Ile Ser Ala Lys Tyr Ser Glu Ser Pro Ser Thr Val
 195 200 205

tcc tgaggtaggc atttgggggc caaggcaaca ggttgggaga attgagaaca 676
 Ser

atggaggaag agtatcttag ggtgccttat ggtggacact cacaaacttg gccatataga 736
 tgtatgtact accagatgaa cagccagcca gattcacaca cgcacatgtt taaacgtgta 796
 aacgtgtgca caactgcaca cacaacctga gaaaccttca ggaggatttg tgggtgtgact 856
 ttgcagtgac atgtagcgat ggctagttga aggaatctcc ctcattgtctt agtgggtcatg 916
 gccacttccc caccctcgcc catctgtgtt cctgcctggc cttgggtggtg cttccgtgtg 976
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 aggtttgagc caccctcccc ttgctagaga gaaggcgn 1074

<210> 8
 <211> 208
 <212> PRT
 <213> Mus musculus

<400> 8

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 Tyr Ile Cys Leu Gly Ala Asn Thr Met Gly Tyr Ser Phe Arg Ser Ala
 35 40 45
 Phe Leu Thr Val Leu Pro Asp Pro Lys Pro Pro Gly Pro Pro Met Ala
 50 55 60
 Ser Ser Ser Ser Thr Ser Leu Pro Trp Pro Val Val Ile Gly Ile
 65 70 75 80
 Pro Ala Gly Ala Val Phe Ile Leu Gly Thr Val Leu Leu Trp Leu Cys
 85 90 95
 Gln Thr Lys Lys Lys Pro Cys Ala Pro Ala Ser Thr Leu Pro Val Pro
 100 105 110
 Gly His Arg Pro Pro Gly Thr Ser Arg Glu Arg Ser Gly Asp Lys Asp
 115 120 125
 Leu Pro Ser Leu Ala Val Gly Ile Cys Glu Glu His Gly Ser Ala Met
 130 135 140
 Ala Pro Gln His Ile Leu Ala Ser Gly Ser Thr Ala Gly Pro Lys Leu
 145 150 155 160
 Tyr Pro Lys Leu Tyr Thr Asp Val His Thr His Thr His Thr
 165 170 175
 Cys Thr His Thr Leu Ser Cys Trp Arg Ala Arg Phe Ile Asn Thr Ser
 180 185 190
 Met Ser Thr Ile Ser Ala Lys Tyr Ser Glu Ser Pro Ser Thr Val Ser
 195 200 205

<210> 9
 <211> 624
 <212> DNA

<213> Mus musculus

<400> 9

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atggggctaca	gtttcogtag	cgcttctctc	actgtattac	cagaccccaa	acctccaggg	180
cctcctatgg	cttcttcato	gtcatccaca	agcctgccat	ggcctgtggt	gacgggcatc	240
ccagctgggtg	ctgtcttcat	cctaggcact	gtgctgctct	ggctttgcca	gaccaagaag	300
aagccatgtg	ccccagcato	tacacttcct	gtgcctgggc	atcgtccccc	agggacatcc	360
cgagaacgca	gtggtgacaa	ggacctgccc	tcattggctg	tgggcatatg	tgaggagcat	420
ggatccgcca	tggcccccca	gcacatcctg	gcctctggct	caactgctgg	ccccaaagctg	480
tacccaagc	tatacacaga	tytgacacaa	cacacacata	cacacacctg	cactcacacg	540
ctctcatgtt	ggagggaag	gttcatcaac	accagcatgt	ccactatcag	tgctaataac	600
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<210> 10

<211> 1423

<212> DNA

<213> Homo sapiens

<220>

<221> CDS

<222> (31)...(444)

<400> 10

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			1				5				
aaa	tat	ctc	tgg	aga	agc	cct	cac	tcc	aaa	ggc	102
Lys	Tyr	Leu	Trp	Arg	Ser	Pro	His	Ser	Lys	Gly	
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tgg	tgg	ctg	ctt	ctc	tgg	gga	gtc	ctc	cag	gct	150
Trp	Trp	Leu	Leu	Leu	Trp	Gly	Val	Leu	Gln	Ala	
25					30				35		40
tcc	gtc	ctc	ttg	gcc	caa	gag	cta	ccc	cag	cag	198
Ser	Val	Leu	Leu	Ala	Gln	Glu	Leu	Pro	Gln	Gln	
				45				50			55
tac	cca	gag	ccg	tat	ggc	aaa	ggc	caa	gag	agc	246
Tyr	Pro	Glu	Pro	Tyr	Gly	Lys	Gly	Gln	Glu	Ser	
			60				65				70
gct	cca	gag	ggc	ttt	gct	gtg	agg	ctc	gtc	ttc	294
Ala	Pro	Glu	Gly	Phe	Ala	Val	Arg	Leu	Val	Phe	
		75					80				85
gag	ccg	tcc	cag	gac	tgt	gca	ggg	gac	tct	gtc	342
Glu	Pro	Ser	Gln	Asp	Cys	Ala	Gly	Asp	Ser	Val	
		90				95				100	
tgg	ggg	ggg	tcc	cgc	cag	gac	tgt	ggc	cag	gga	390
Trp	Gly	Gly	Ser	Arg	Gln	Asp	Cys	Gly	Gln	Gly	
105					110				115		120
ggg	aag	tgg	cgg	tgc	cct	gaa	tcc	ccc	atc	tgg	438

Gly Lys Trp Arg Cys Pro Glu Ser Pro Ile Trp Arg Arg Asp Glu Phe
125 130 135

tcc atg taggggcagt cgggcttgcc ttaccgggga gcagtgggtg accccaggac 494
Ser Met

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caagcaggcc ctgogtttgg aaggcttatg aatggacaca caaatcttgc aaatctatgg 614
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tgctgtctct atcaggtgag gaagctggac acaataata acaaaagatt aagtcaccgt 734
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caacatagca agaccocatc tcaaaaataa gtaataata aataaaaaata aaaagagcac 974
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caaaaaaaaa aaaaaaaaaag ggcggccgc 1423

<210> 11
<211> 138
<212> PRT
<213> Homo sapiens

<400> 11
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20 25 30
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35 40 45
Pro Gln Gln Leu Thr Ser Pro Gly Tyr Pro Glu Pro Tyr Gly Lys Gly
50 55 60
Gln Glu Ser Ser Thr Asp Ile Lys Ala Pro Glu Gly Phe Ala Val Arg
65 70 75 80
Leu Val Phe Gln Asp Phe Asp Leu Glu Pro Ser Gln Asp Cys Ala Gly
85 90 95
Asp Ser Val Thr Val Ser Trp Gly Trp Gly Gly Ser Arg Gln Asp Cys
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Gly Gln Gly Asp Ser Arg Gly Cys Gly Lys Trp Arg Cys Pro Glu Ser
115 120 125
Pro Ile Trp Arg Arg Asp Glu Phe Ser Met
130 135

<210> 12
<211> 414
<212> DNA
<213> Homo sapiens

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tccgtctctc tggcccaaga gctacccag cagctgacat ccccgggta cccagagccg 180

tatggcaaag gccaaagagag cagcacggac atcaaggctc cagagggctt tgctgtgagg	240
ctcgtcttcc aggacttcca cctggagccg tcccaggact gtgcagggga ctctgtcaca	300
gtgagctggg gatggggggg gtcccgccag gactgtggcc agggagattc ccggggttgt	360
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<210> 13
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<220>
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aggettata tctgaacgc tgggatcccc caggacattc cctggcccc agggcccagg	180
tcccaggccc cagggtgag ctgtgggcag gccccacctg gcctctgca atg tca ccg	238
	Met Ser Pro
	1

cct ctg tgt ccc ctc ctt ctc ctg gct gtg ggc ctg cgg ctg gct gga	286
Pro Leu Cys Pro Leu Leu Leu Leu Ala Val Gly Leu Arg Leu Ala Gly	
5 10 15	

act ctc aac ccc agt gat ccc aat acc tgc agc ttc tgg gaa agc ttc	334
Thr Leu Asn Pro Ser Asp Pro Asn Thr Cys Ser Phe Trp Glu Ser Phe	
20 25 30 35	

act acc acc acc aag gag tcc cac tcc cgc ccc ttc agc ctg ctc ccc	382
Thr Thr Thr Thr Lys Glu Ser His Ser Arg Pro Phe Ser Leu Leu Pro	
40 45 50	

tca gag ccc tgc gag cgg ccc tgg gag ggc ccc cat act tgc ccc agc	430
Ser Glu Pro Cys Glu Arg Pro Trp Glu Gly Pro His Thr Cys Pro Ser	
55 60 65	

cca caa act cag agg aaa ctc ctg gct tct agg gat tca ttc tgc atg	478
Pro Gln Thr Gln Arg Lys Leu Leu Ala Ser Arg Asp Ser Phe Cys Met	
70 75 80	

gtc tgt gtc ggg gct gga gtg cag tgg cga gat cgt agt gca ctg caa	526
Val Cys Val Gly Ala Gly Val Gln Trp Arg Asp Arg Ser Ala Leu Gln	
85 90 95	

cct caa aca ggg aat gcg ctt tct atg cgc cct cag ccc aga gtg ttg	574
Pro Gln Thr Gly Asn Ala Leu Ser Met Arg Pro Gln Pro Arg Val Leu	
100 105 110 115	

agt ggt gcc cct tcc ctg gcc tcc cct ggc cac act gtg gtg gtg aag	622
Ser Gly Ala Pro Ser Leu Ala Ser Pro Gly His Thr Val Val Val Lys	
120 125 130	

acg gac cac cgc cag cgc ctg cag tgc tgc cat ggc ttc tat gag agc	670
Thr Asp His Arg Gln Arg Leu Gln Cys Cys His Gly Phe Tyr Glu Ser	
135 140 145	

agg ggg ttc tgt gtc ccg ctc tgt gcc cag gag tgt gtc cat ggc cgt	718
Arg Gly Phe Cys Val Pro Leu Cys Ala Gln Glu Cys Val His Gly Arg	
150 155 160	
tgt gtg gca ccc aat cag tgc caa tgt gtg cca ggc tgg cgg ggc gac	766
Cys Val Ala Pro Asn Gln Cys Gln Cys Val Pro Gly Trp Arg Gly Asp	
165 170 175	
gac tgt tcc agt gcc ccg aac tgc ctt cag ccc tgt acc cct ggc tac	814
Asp Cys Ser Ser Ala Pro Asn Cys Leu Gln Pro Cys Thr Pro Gly Tyr	
180 185 190 195	
tat ggc cct gcc tgc cag ttc cgc tgc cag tgc cat ggg gca ccc tgc	862
Tyr Gly Pro Ala Cys Gln Phe Arg Cys Gln Cys His Gly Ala Pro Cys	
200 205 210	
gat ccc cag act gga gcc tgc ttc tgc ccc gca gag aga act ggg ccc	910
Asp Pro Gln Thr Gly Ala Cys Phe Cys Pro Ala Glu Arg Thr Gly Pro	
215 220 225	
agc tgt gac gtg tcc tgt tcc cag ggc act tct ggc ttc ttc tgc ccc	958
Ser Cys Asp Val Ser Cys Ser Gln Gly Thr Ser Gly Phe Cys Pro	
230 235 240	
agc acc cat cct tgc caa aat gga ggt gtc ttc caa acc cca cag ggc	1006
Ser Thr His Pro Cys Gln Asn Gly Gly Val Phe Gln Thr Pro Gln Gly	
245 250 255	
tcc tgc agc tgc ccc cct ggc tgg atg ggc acc atc tgc tcc ctg ccc	1054
Ser Cys Ser Cys Pro Pro Gly Trp Met Gly Thr Ile Cys Ser Leu Pro	
260 265 270 275	
tgc cca gag ggc ttt cac gga ccc aac tgc tcc cag gaa tgt cgc tgc	1102
Cys Pro Glu Gly Phe His Gly Pro Asn Cys Ser Gln Glu Cys Arg Cys	
280 285 290	
cac aac ggc ggc ctc tgt gac cga ttc act ggg cag tgc cgc tgc gct	1150
His Asn Gly Gly Leu Cys Asp Arg Phe Thr Gly Gln Cys Arg Cys Ala	
295 300 305	
ccg ggt tac act ggg gat cgg tgc cgg gag gag tgc ccg gtg ggc cgc	1198
Pro Gly Tyr Thr Gly Asp Arg Cys Arg Glu Glu Cys Pro Val Gly Arg	
310 315 320	
ttt ggg cag gac tgt gct gag acg tgc gac tgc gcc ccg gac gcc cgt	1246
Phe Gly Gln Asp Cys Ala Glu Thr Cys Asp Cys Ala Pro Asp Ala Arg	
325 330 335	
tgc ttc ccg gcc aac ggc gca tgt ctg tgc gaa cac ggc ttc act ggg	1294
Cys Phe Pro Ala Asn Gly Ala Cys Leu Cys Glu His Gly Phe Thr Gly	
340 345 350 355	
gac cgc tgc acg gat cgc ctc tgc ccc gac ggc ttc tac ggt ctc agc	1342
Asp Arg Cys Thr Asp Arg Leu Cys Pro Asp Gly Phe Tyr Gly Leu Ser	
360 365 370	
tgc cag gcc ccc tgc acc tgc gac cgg gag cac agc ctc agc tgc cac	1390
Cys Gln Ala Pro Cys Thr Cys Asp Arg Glu His Ser Leu Ser Cys His	

375	380	385	
ccg atg aac ggg gag tgc tcc tgc ctg ccg ggc tgg gcg ggc ctc cac Pro Met Asn Gly Glu Cys Ser Cys Leu Pro Gly Trp Ala Gly Leu His 390 395 400			1438
tgc aac gag agc tgc ccg cag gac acg cat ggg cca ggg tgc cag gag Cys Asn Glu Ser Cys Pro Gln Asp Thr His Gly Pro Gly Cys Gln Glu 405 410 415			1486
cac tgt ctc tgc ctg cac ggt ggc gtc tgc cag gct acc agc ggc ctc His Cys Leu Cys Leu His Gly Gly Val Cys Gln Ala Thr Ser Gly Leu 420 425 430 435			1534
tgt cag tgc gcg ccg ggt tac acg ggc cct cac tgt gct agt ctt tgt Cys Gln Cys Ala Pro Gly Tyr Thr Gly Pro His Cys Ala Ser Leu Cys 440 445 450			1582
cct cct gac acc tac ggt gtc aac tgt tct gca cgc tgc tca tgt gaa Pro Pro Asp Thr Tyr Gly Val Asn Cys Ser Ala Arg Cys Ser Cys Glu 455 460 465			1630
aat gcc atc gcc tgc tca ccc atc gac ggc gag tgc gtc tgc aag gaa Asn Ala Ile Ala Cys Ser Pro Ile Asp Gly Glu Cys Val Cys Lys Glu 470 475 480			1678
ggt tgg cag cgt ggt aac tgc tct gtg ccc tgc cca ccc gga acc tgg Gly Trp Gln Arg Gly Asn Cys Ser Val Pro Cys Pro Pro Gly Thr Trp 485 490 495			1726
ggc ttc agt tgc aat gcc agc tgc cag tgt gcc cat gag gca gtc tgc Gly Phe Ser Cys Asn Ala Ser Cys Gln Cys Ala His Glu Ala Val Cys 500 505 510 515			1774
agc ccc caa act gga gcc tgt acc tgc acc cct ggg tgg cat ggg gcc Ser Pro Gln Thr Gly Ala Cys Thr Cys Thr Pro Gly Trp His Gly Ala 520 525 530			1822
cac tgc cag ctg ccc tgt ccg aag ggg cag ttt gga gaa ggt tgt gcc His Cys Gln Leu Pro Cys Pro Lys Gly Gln Phe Gly Glu Gly Cys Ala 535 540 545			1870
agt cgc tgt gac tgt gac cac tct gat ggc tgt gac cct gtt cat gga Ser Arg Cys Asp Cys Asp His Ser Asp Gly Cys Asp Pro Val His Gly 550 555 560			1918
cgc tgt cag tgc cag gct ggc tgg atg ggt gcc cgc tgc cac ctg tcc Arg Cys Gln Cys Gln Ala Gly Trp Met Gly Ala Arg Cys His Leu Ser 565 570 575			1966
tgc cct gag ggc tta tgg gga gtc aac tgt agc aac acc tgc acc tgc Cys Pro Glu Gly Leu Trp Gly Val Asn Cys Ser Asn Thr Cys Thr Cys 580 585 590 595			2014
aag aat ggg ggc acc tgt ctc cct gag aat ggc aac tgc gtg tgt gca Lys Asn Gly Gly Thr Cys Leu Pro Glu Asn Gly Asn Cys Val Cys Ala 600 605 610			2062

ccc gga ttc cgg ggc ccc tcc tgc cag aga tcc tgt cag cct ggc cgc Pro Gly Phe Arg Gly Pro Ser Cys Gln Arg Ser Cys Gln Pro Gly Arg 615 620 625	2110
tat ggc aaa cgc tgt gtg ccc tgc aag tgc gct aac cac tcc ttc tgc Tyr Gly Lys Arg Cys Val Pro Cys Lys Cys Ala Asn His Ser Phe Cys 630 635 640	2158
cac ccc tcg aac ggg acc tgc tac tgc ctg gct ggc tgg aca ggc ccc His Pro Ser Asn Gly Thr Cys Tyr Cys Leu Ala Gly Trp Thr Gly Pro 645 650 655	2206
gac tgc tcc cag cca tgc cct cca gga cac tgg gga gaa aac tgt gcc Asp Cys Ser Gln Pro Cys Pro Pro Gly His Trp Gly Glu Asn Cys Ala 660 665 670 675	2254
cag acc tgc caa tgt cac cat ggt ggg acc tgc cat ccc cag gat ggg Gln Thr Cys Gln Cys His His Gly Gly Thr Cys His Pro Gln Asp Gly 680 685 690	2302
agc tgt atc tgc ccc cta ggc tgg act gga cac cac tgc tta gaa ggc Ser Cys Ile Cys Pro Leu Gly Trp Thr Gly His His Cys Leu Glu Gly 695 700 705	2350
tgc cct ctg ggg aca ttt ggt gct aac tgc tcc cag cca tgc cag tgt Cys Pro Leu Gly Thr Phe Gly Ala Asn Cys Ser Gln Pro Cys Gln Cys 710 715 720	2398
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cca ggg cac agt ggt gca cct tgc agg att gga atc cag gag ccc ttt Pro Gly His Ser Gly Ala Pro Cys Arg Ile Gly Ile Gln Glu Pro Phe 740 745 750 755	2494
act gtg atg ccg acc act cca gta gcg tat aac tcg ctg ggt gca gtg Thr Val Met Pro Thr Thr Pro Val Ala Tyr Asn Ser Leu Gly Ala Val 760 765 770	2542
att ggc att gca gtg ctg ggg tcc ctt gtg gta gcc ctg gtg gca ctg Ile Gly Ile Ala Val Leu Gly Ser Leu Val Val Ala Leu Val Ala Leu 775 780 785	2590
ttc att ggc tat cgg cac tgg caa aaa ggc aag gag cac cac cac ctg Phe Ile Gly Tyr Arg His Trp Gln Lys Gly Lys Glu His His His Leu 790 795 800	2638
gct gtg gct tac agc agc ggg cgc ctg gac ggc tcc gag tat gtc atg Ala Val Ala Tyr Ser Ser Gly Arg Leu Asp Gly Ser Glu Tyr Val Met 805 810 815	2686
cca gat gtc cct ccg agc tac agt cac tac tac tcc aac ccc agc tac Pro Asp Val Pro Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr 820 825 830 835	2734
cac acc ctg tcg cag tgc tcc cca aac ccc cca ccc cct aac aag gtt His Thr Leu Ser Gln Cys Ser Pro Asn Pro Pro Pro Pro Asn Lys Val	2782

840	845	850	
cca ggc cgc ctc ttt gcc agc ctg cag aac cct gag cgg cca ggt ggg			2830
Pro Gly Pro Leu Phe Ala Ser Leu Gln Asn Pro Glu Arg Pro Gly Gly	855	860 865	
gcc caa ggg cat gat aac cac acc acc ctg cct gct gac tgg aag cac			2878
Ala Gln Gly His Asp Asn His Thr Thr Leu Pro Ala Asp Trp Lys His	870	875 880	
cgc cgg gag ccc cct cca ggg cct ctg gac agg ggg agc agc cgc ctg			2926
Arg Arg Glu Pro Pro Pro Gly Pro Leu Asp Arg Gly Ser Ser Arg Leu	885	890 895	
gac cga agc tac agc tat agc tac agc aat ggc cca ggc cca ttc tac			2974
Asp Arg Ser Tyr Ser Tyr Ser Tyr Ser Asn Gly Pro Gly Pro Phe Tyr	900	905 910 915	
gat aaa ggg ctc atc tct gaa gag gag ctc ggg gcc agt gtg gct tcc			3022
Asp Lys Gly Leu Ile Ser Glu Glu Glu Leu Gly Ala Ser Val Ala Ser	920	925 930	
ctg agc agt gag aac cca tat gcc acc atc cgg gac ctg ccc agc ttg			3070
Leu Ser Ser Glu Asn Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu	935	940 945	
cca ggg ggc ccc cgg gag agc agc tac atg gag atg aaa ggc cct ccc			3118
Pro Gly Gly Pro Arg Glu Ser Ser Tyr Met Glu Met Lys Gly Pro Pro	950	955 960	
tca gga tct gcc ccc agg cag cct cct cag ttt tgg gac agc cag agg			3166
Ser Gly Ser Ala Pro Arg Gln Pro Pro Gln Phe Trp Asp Ser Gln Arg	965	970 975	
cgg cgg caa ccc cag cca cag aga gac agt ggc acc tac gag cag ccc			3214
Arg Arg Gln Pro Gln Pro Gln Arg Asp Ser Gly Thr Tyr Glu Gln Pro	980	985 990 995	
agc ccc ctg atc cat gac cga gac tct gtg ggc tcc cag ccc cct ctg			3262
Ser Pro Leu Ile His Asp Arg Asp Ser Val Gly Ser Gln Pro Pro Leu	1000	1005 1010	
cct ccg ggc cta ccc ccc ggc cac tat gac tca ccc aag aac agc cac			3310
Pro Pro Gly Leu Pro Pro Gly His Tyr Asp Ser Pro Lys Asn Ser His	1015	1020 1025	
atc cct gga cat tat gac ttg cct cca gta cgg cat ccc cca tca cct			3358
Ile Pro Gly His Tyr Asp Leu Pro Pro Val Arg His Pro Pro Ser Pro	1030	1035 1040	
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Pro Leu Arg Arg Gln Asp Arg	1045	1050	
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ctctacattg	cattttggta	attgcttgca	atatttcaag	cattttcatt	gttattatat	4909
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<210> 14
 <211> 1050
 <212> PRT
 <213> Homo sapiens

<400> 14

Met	Ser	Pro	Pro	Cys	Pro	Leu	Leu	Leu	Leu	Ala	Val	Gly	Leu	Arg
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Leu	Ala	Gly	Thr	Leu	Asn	Pro	Ser	Asp	Pro	Asn	Thr	Cys	Ser	Phe
			20				25					30		
Glu	Ser	Phe	Thr	Thr	Thr	Lys	Glu	Ser	His	Ser	Arg	Pro	Phe	Ser
			35			40					45			
Leu	Leu	Pro	Ser	Glu	Pro	Cys	Glu	Arg	Pro	Trp	Glu	Gly	Pro	His
			50			55					60			
Cys	Pro	Ser	Pro	Gln	Thr	Gln	Arg	Lys	Leu	Leu	Ala	Ser	Arg	Asp
65				70					75					80
Phe	Cys	Met	Val	Cys	Val	Gly	Ala	Gly	Val	Gln	Trp	Arg	Asp	Arg
			85						90					95
Ala	Leu	Gln	Pro	Gln	Thr	Gly	Asn	Ala	Leu	Ser	Met	Arg	Pro	Gln
			100					105					110	
Arg	Val	Leu	Ser	Gly	Ala	Pro	Ser	Leu	Ala	Ser	Pro	Gly	His	Thr
			115					120				125		
Val	Val	Lys	Thr	Asp	His	Arg	Gln	Arg	Leu	Gln	Cys	Cys	His	Gly
			130				135				140			
Tyr	Glu	Ser	Arg	Gly	Phe	Cys	Val	Pro	Leu	Cys	Ala	Gln	Glu	Cys
145				150					155					160
His	Gly	Arg	Cys	Val	Ala	Pro	Asn	Gln	Cys	Gln	Cys	Val	Pro	Gly
			165					170					175	
Arg	Gly	Asp	Asp	Cys	Ser	Ser	Ala	Pro	Asn	Cys	Leu	Gln	Pro	Cys
			180					185					190	
Pro	Gly	Tyr	Tyr	Gly	Pro	Ala	Cys	Gln	Phe	Arg	Cys	Gln	Cys	His
			195				200					205		
Ala	Pro	Cys	Asp	Pro	Gln	Thr	Gly	Ala	Cys	Phe	Cys	Pro	Ala	Glu
														Arg

210	215	220
Thr Gly Pro Ser Cys Asp	Val Ser Cys Ser Gln Gly Thr Ser Gly Phe	
225	230	235
Phe Cys Pro Ser Thr His	Pro Cys Gln Asn Gly Gly Val Phe Gln Thr	240
245	250	255
Pro Gln Gly Ser Cys Ser Cys	Pro Pro Gly Trp Met Gly Thr Ile Cys	
260	265	270
Ser Leu Pro Cys Pro Glu Gly	Phe His Gly Pro Asn Cys Ser Gln Glu	
275	280	285
Cys Arg Cys His Asn Gly	Gly Leu Cys Asp Arg Phe Thr Gly Gln Cys	
290	295	300
Arg Cys Ala Pro Gly Tyr Thr	Gly Asp Arg Cys Arg Glu Glu Cys Pro	
305	310	315
Val Gly Arg Phe Gly Gln Asp	Cys Ala Glu Thr Cys Asp Cys Ala Pro	320
325	330	335
Asp Ala Arg Cys Phe Pro Ala	Asn Gly Ala Cys Leu Cys Glu His Gly	
340	345	350
Phe Thr Gly Asp Arg Cys Thr	Asp Arg Leu Cys Pro Asp Gly Phe Tyr	
355	360	365
Gly Leu Ser Cys Gln Ala Pro	Cys Thr Cys Asp Arg Glu His Ser Leu	
370	375	380
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Gly Leu His Cys Asn Glu Ser	Cys Pro Gln Asp Thr His Gly Pro Gly	400
405	410	415
Cys Gln Glu His Cys Leu Cys	Leu His Gly Gly Val Cys Gln Ala Thr	
420	425	430
Ser Gly Leu Cys Gln Cys Ala	Pro Gly Tyr Thr Gly Pro His Cys Ala	
435	440	445
Ser Leu Cys Pro Pro Asp Thr	Tyr Gly Val Asn Cys Ser Ala Arg Cys	
450	455	460
Ser Cys Glu Asn Ala Ile Ala	Cys Ser Pro Ile Asp Gly Glu Cys Val	
465	470	475
Cys Lys Glu Gly Trp Gln Arg	Gly Asn Cys Ser Val Pro Cys Pro Pro	
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Gly Thr Trp Gly Phe Ser Cys	Asn Ala Ser Cys Gln Cys Ala His Glu	
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Ala Val Cys Ser Pro Gln Thr	Gly Ala Cys Thr Cys Thr Pro Gly Trp	
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His Gly Ala His Cys Gln Leu	Pro Cys Pro Lys Gly Gln Phe Gly Glu	
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Gly Cys Ala Ser Arg Cys Asp	Cys Asp His Ser Asp Gly Cys Asp Pro	
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Val His Gly Arg Cys Gln Cys	Gln Ala Gly Trp Met Gly Ala Arg Cys	
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His Leu Ser Cys Pro Glu Gly	Leu Trp Gly Val Asn Cys Ser Asn Thr	
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Cys Thr Cys Lys Asn Gly Gly	Thr Cys Leu Pro Glu Asn Gly Asn Cys	
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Val Cys Ala Pro Gly Phe Arg	Gly Pro Ser Cys Gln Arg Ser Cys Gln	
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Ser Phe Cys His Pro Ser Asn	Gly Thr Cys Tyr Cys Leu Ala Gly Trp	
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Thr Gly Pro Asp Cys Ser Gln	Pro Cys Pro Pro Gly His Trp Gly Glu	
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Asn Cys Ala Gln Thr Cys Gln	Cys His His Gly Gly Thr Cys His Pro	

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705	Cys Gln Cys Gly Pro Gly	710	Glu Lys Cys His Pro	715	Glu Thr Gly Ala Cys
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	Glu Pro Phe Thr Val Met		Pro Thr Thr Pro Val		Ala Tyr Asn Ser Leu
		755		760	
	Gly Ala Val Ile Gly Ile		Ala Val Leu Gly Ser		Leu Val Val Ala Leu
		770		775	
	Val Ala Leu Phe Ile Gly		Tyr Arg His Trp Gln		Lys Gly Lys Glu His
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	His His Leu Ala Val Ala		Tyr Ser Ser Gly Arg		Leu Asp Gly Ser Glu
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		820		825	
	Pro Ser Tyr His Thr Leu		Ser Gln Cys Ser Pro		Asn Pro Pro Pro Pro
		835		840	
	Asn Lys Val Pro Gly Pro		Leu Phe Ala Ser Leu		Gln Asn Pro Glu Arg
		850		855	
	Pro Gly Gly Ala Gln Gly		His Asp Asn His Thr		Thr Leu Pro Ala Asp
		865		870	
	Trp Lys His Arg Arg Glu		Pro Pro Pro Gly Pro		Leu Asp Arg Gly Ser
		885		890	
	Ser Arg Leu Asp Arg Ser		Tyr Ser Tyr Ser Tyr		Ser Asn Gly Pro Gly
		900		905	
	Pro Phe Tyr Asp Lys Gly		Leu Ile Ser Glu Glu		Glu Glu Leu Gly Ala Ser
		915		920	
	Val Ala Ser Leu Ser Ser		Glu Asn Pro Tyr Ala		Thr Ile Arg Asp Leu
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	Ser Gln Arg Arg Arg Gln		Pro Gln Pro Gln Arg		Asp Ser Gly Thr Tyr
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	Glu Gln Pro Ser Pro Leu		Ile His Asp Arg Asp		Ser Val Gly Ser Gln
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	Asn Ser His Ile Pro Gly		His Tyr Asp Leu Pro		Pro Pro Val Arg His Pro
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 <212> DNA
 <213> Mus musculus

<220>
 <221> CDS
 <222> (2)...(1492)

<400> 16

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gct	ggg	tgg	atg	ggc	aca	cgc	tgc	cac	ctg	cct	tgc	ccg	gag	ggc	ttt	97	
Ala	Gly	Trp	Met	Gly	Thr	Arg	Cys	His	Leu	Pro	Cys	Pro	Glu	Gly	Phe		
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Trp	Gly	Ala	Asn	Cys	Ser	Asn	Thr	Cys	Thr	Cys	Lys	Asn	Gly	Gly	Thr		
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Cys	Val	Ser	Glu	Asn	Gly	Asn	Cys	Val	Cys	Ala	Pro	Gly	Phe	Arg	Gly		
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Pro	Ser	Cys	Gln	Arg	Pro	Cys	Pro	Pro	Gly	Arg	Tyr	Gly	Lys	Arg	Cys		
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Val	Gln	Cys	Lys	Cys	Asn	Asn	Asn	His	Ser	Ser	Cys	His	Pro	Ser	Asp		
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Gly	Thr	Cys	Ser	Cys	Leu	Ala	Gly	Trp	Thr	Gly	Pro	Asp	Cys	Ser	Glu		
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gca	tgt	ccc	cca	ggc	cac	tgg	gga	ctc	aaa	tgc	tcc	caa	ctc	tgc	cag	385	
Ala	Cys	Pro	Pro	Gly	His	Trp	Gly	Leu	Lys	Cys	Ser	Gln	Leu	Cys	Gln		
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Cys	His	His	Gly	Gly	Thr	Cys	His	Pro	Gln	Asp	Gly	Ser	Cys	Ile	Cys		
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acg	cca	ggc	tgg	act	gga	ccc	aac	tgc	ttg	gaa	ggc	tgc	cca	cca	aga	481	
Thr	Pro	Gly	Trp	Thr	Gly	Pro	Asn	Cys	Leu	Glu	Gly	Cys	Pro	Pro	Arg		
	145				150				155						160		
atg	ttt	ggg	gtc	aac	tgc	tcc	cag	cta	tgt	cag	tgt	gat	ctc	gga	gag	529	
Met	Phe	Gly	Val	Asn	Cys	Ser	Gln	Leu	Cys	Gln	Cys	Asp	Leu	Gly	Glu		
			165					170				175					
atg	tgc	cac	cca	gag	act	ggg	gct	tgt	gtc	tgt	ccc	cca	gga	cac	agt	577	
Met	Cys	His	Pro	Glu	Thr	Gly	Ala	Cys	Val	Cys	Pro	Pro	Gly	His	Ser		
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ggg	gca	gac	tgc	aaa	atg	gga	agc	cag	gag	tcc	ttc	acc	ata	atg	ccc	625	
Gly	Ala	Asp	Cys	Lys	Met	Gly	Ser	Gln	Glu	Ser	Phe	Thr	Ile	Met	Pro		
		195				200						205					
acc	tct	ccc	gtg	acc	cat	aac	tca	ctg	ggg	gca	gtg	att	ggc	att	gca	673	
Thr	Ser	Pro	Val	Thr	His	Asn	Ser	Leu	Gly	Ala	Val	Ile	Gly	Ile	Ala		
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gta	ctg	gga	acc	ctc	gtg	gtg	gcc	ctg	ata	gca	ctg	ttc	att	ggc	tac	721	

Val Leu Gly Thr Leu Val Val Ala Leu Ile Ala Leu Phe Ile Gly Tyr	
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cgc cag tgg caa aag ggc aag gaa cat gag cac ttg gca gtg gct tac	769
Arg Gln Trp Gln Lys Gly Lys Glu His Glu His Leu Ala Val Ala Tyr	
245 250 255	
agc act ggg cgg ctg gat ggc tct gat tac gtc atg cca gat gtc tct	817
Ser Thr Gly Arg Leu Asp Gly Ser Asn Tyr Val Met Pro Asp Val Ser	
260 265 270	
ccg agc tat agt cac tac tac tcc aac ccc agc tac cac aca ctg tct	865
Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser	
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Gln Cys Ser Pro Asn Pro Pro Pro Asn Lys Val Pro Gly Ser Gln	
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Leu Phe Val Ser Ser Gln Ala Pro Glu Arg Pro Ser Arg Ala His Gly	
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cgt gag aac cat acc aca ctg ccc gct gac tgg aag cac cgc cgg gag	1009
Arg Glu Asn His Thr Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu	
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ccc cat gac aga ggc gcc agc cac ctg gac cga agc tat agc tgt agc	1057
Pro His Asp Arg Gly Ala Ser His Leu Asp Arg Ser Tyr Ser Cys Ser	
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Tyr Ser His Arg Asn Gly Pro Gly Pro Phe Cys His Lys Gly Pro Ile	
355 360 365	
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Ser Glu Glu Gly Leu Gly Ala Ser Val Met Ser Leu Ser Ser Glu Asn	
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Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu Pro Gly Glu Pro Arg	
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Glu Ser Gly Tyr Val Glu Met Lys Gly Pro Pro Ser Val Ser Pro Pro	
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Arg Gln Ser Leu His Leu Arg Asp Arg Gln Gln Arg Gln Leu Gln Pro	
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Gln Arg Asp Ser Gly Thr Tyr Glu Gln Pro Ser Pro Leu Ser His Asn	
435 440 445	
gaa gag tct ttg ggc tcc acg ccc ccg ctt cct cca ggc ctg cct cct	1393
Glu Glu Ser Leu Gly Ser Thr Pro Pro Leu Pro Gly Leu Pro Pro	
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 Gly His Tyr Asp Ser Pro Lys Asn Ser His Ile Pro Gly His Tyr Asp
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 Leu Pro Pro Val Arg His Pro Pro Ser Pro Pro Ser Arg Arg Gln Asp
 485 490 495

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<210> 17
 <211> 497
 <212> PRT
 <213> Mus musculus

<400> 17
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 35 40 45
 Cys Val Ser Glu Asn Gly Asn Cys Val Cys Ala Pro Gly Phe Arg Gly
 50 55 60
 Pro Ser Cys Gln Arg Pro Cys Pro Pro Gly Arg Tyr Gly Lys Arg Cys
 65 70 75 80
 Val Gln Cys Lys Cys Asn Asn Asn His Ser Ser Cys His Pro Ser Asp
 85 90 95
 Gly Thr Cys Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys Ser Glu
 100 105 110
 Ala Cys Pro Pro Gly His Trp Gly Leu Lys Cys Ser Gln Leu Cys Gln
 115 120 125
 Cys His His Gly Gly Thr Cys His Pro Gln Asp Gly Ser Cys Ile Cys
 130 135 140
 Thr Pro Gly Trp Thr Gly Pro Asn Cys Leu Glu Gly Cys Pro Pro Arg
 145 150 155 160

Met Phe Gly Val Asn Cys Ser Gln Leu Cys Gln Cys Asp Leu Gly Glu
165 170 175
Met Cys His Pro Glu Thr Gly Ala Cys Val Cys Pro Pro Gly His Ser
180 185 190
Gly Ala Asp Cys Lys Met Gly Ser Gln Glu Ser Phe Thr Ile Met Pro
195 200 205
Thr Ser Pro Val Thr His Asn Ser Leu Gly Ala Val Ile Gly Ile Ala
210 215 220
Val Leu Gly Thr Leu Val Val Ala Leu Ile Ala Leu Phe Ile Gly Tyr
225 230 235 240
Arg Gln Trp Gln Lys Gly Lys Glu His Glu His Leu Ala Val Ala Tyr
245 250 255
Ser Thr Gly Arg Leu Asp Gly Ser Asp Tyr Val Met Pro Asp Val Ser
260 265 270
Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser
275 280 285
Gln Cys Ser Pro Asn Pro Pro Pro Asn Lys Val Pro Gly Ser Gln
290 295 300
Leu Phe Val Ser Ser Gln Ala Pro Glu Arg Pro Ser Arg Ala His Gly
305 310 315 320
Arg Glu Asn His Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu
325 330 335
Pro His Asp Arg Gly Ala Ser His Leu Asp Arg Ser Tyr Ser Cys Ser
340 345 350
Tyr Ser His Arg Asn Gly Pro Gly Pro Phe Cys His Lys Gly Pro Ile
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Ser Glu Glu Gly Leu Gly Ala Ser Val Met Ser Leu Ser Ser Glu Asn
370 375 380
Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu Pro Gly Glu Pro Arg
385 390 395 400
Glu Ser Gly Tyr Val Glu Met Lys Gly Pro Pro Ser Val Ser Pro Pro
405 410 415
Arg Gln Ser Leu His Leu Arg Asp Arg Gln Gln Arg Gln Leu Gln Pro
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Gln Arg Asp Ser Gly Thr Tyr Glu Gln Pro Ser Pro Leu Ser His Asn
435 440 445
Glu Glu Ser Leu Gly Ser Thr Pro Pro Leu Pro Pro Gly Leu Pro Pro
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485 490 495
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<211> 1491
<212> DNA
<213> Mus musculus

<400> 18
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caggagtcc	tcaccataat	gcccacctct	cccgtgaccc	ataactcact	gggtgcagt	660
attggcattg	cagtactggg	aaccctcgtg	gtggccctga	tagcactgtt	cattggctac	720
cgccagtggc	aaaaggggcaa	ggaacatgag	cacttggcag	tggtttacag	cactgggcgg	780
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tcctgcagct	gcccaccggg	ctgg atg ggt gtc atc tgt tcc ctg cca tgc				951

Met Gly Val Ile Cys Ser Leu Pro Cys
 1 5

cca gag ggt ttc cac gga ccc aac tgt act cag gaa tgt cgt tgc cac	999
Pro Glu Gly Phe His Gly Pro Asn Cys Thr Gln Glu Cys Arg Cys His	
10 15 20 25	

aat ggt ggc ctt tgt gac agg ttt act ggg cag tgc cac tgt gct cct	1047
Asn Gly Gly Leu Cys Asp Arg Phe Thr Gly Gln Cys His Cys Ala Pro	
30 35 40	

ggc tat atc ggg gat cgg tgc cgt gaa gag tgc cct gtg ggc cgc ttc	1095
Gly Tyr Ile Gly Asp Arg Cys Arg Glu Glu Cys Pro Val Gly Arg Phe	

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	Gly Gln Asp Cys Ala Glu Thr	Cys Asp Cys Ala Pro Gly Ala Arg Cys		
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	ttt cct gcc aat ggc gcg tgt ctg tgc gaa cat ggc ttc aca ggc gac		1191	
	Phe Pro Ala Asn Gly Ala Cys Leu Cys Glu His Gly Phe Thr Gly Asp			
	75	80	85	
	cgc tgc act gag cga ctc tgt cca gat ggc cgc tat ggt ctg agc tgc		1239	
	Arg Cys Thr Glu Arg Leu Cys Pro Asp Gly Arg Tyr Gly Leu Ser Cys			
	90	95	100	105
	caa gat ccc tgc acc tgc gac cca gaa cac agt ctc agc tgc cac cca		1287	
	Gln Asp Pro Cys Thr Cys Asp Pro Glu His Ser Leu Ser Cys His Pro			
	110	115	120	
	atg cac ggc gag tgc tcc tgc cag cca ggt tgg gcg ggc ctc cac tgc		1335	
	Met His Gly Glu Cys Ser Cys Gln Pro Gly Trp Ala Gly Leu His Cys			
	125	130	135	
	aac gag agc tgc cct cag gac acg cac gga gcc ggt tgc cag gag cac		1383	
	Asn Glu Ser Cys Pro Gln Asp Thr His Gly Ala Gly Cys Gln Glu His			
	140	145	150	
	tgc ctc tgt ctg cac ggc ggt gtt tgc ctc gcc gac agc ggc ctc tgc		1431	
	Cys Leu Cys Leu His Gly Gly Val Cys Leu Ala Asp Ser Gly Leu Cys			
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	cgg tgt gca cct ggc tac acg gga cct cac tgc gct aat ctt tgt cca		1479	
	Arg Cys Ala Pro Gly Tyr Thr Gly Pro His Cys Ala Asn Leu Cys Pro			
	170	175	180	185
	cct aac act tat ggg atc aac tgt tcc tcc cac tgc tcc tgt gaa aat		1527	
	Pro Asn Thr Tyr Gly Ile Asn Cys Ser Ser His Cys Ser Cys Glu Asn			
	190	195	200	
	gcc att gcc tgc tct cct gtc gac ggc acg tgc atc tgc aag gaa ggt		1575	
	Ala Ile Ala Cys Ser Pro Val Asp Gly Thr Cys Ile Cys Lys Glu Gly			
	205	210	215	
	tgg cag cgt ggt aac tgc tct gtg ccc tgt ccc cct ggc acc tgg ggc		1623	
	Trp Gln Arg Gly Asn Cys Ser Val Pro Cys Pro Pro Gly Thr Trp Gly			
	220	225	230	
	ttc agt tgc aat gcc agt tgc cag tgt gcc cac gag gga gtc tgc agc		1671	
	Phe Ser Cys Asn Ala Ser Cys Gln Cys Ala His Glu Gly Val Cys Ser			
	235	240	245	
	ccc caa act gga gcc tgt act tgc acc cct ggg tgg cgt ggg gtt cac		1719	
	Pro Gln Thr Gly Ala Cys Thr Cys Thr Pro Gly Trp Arg Gly Val His			
	250	255	260	265
	tgc caa ctt ccg tgc ccg aag gga cag ttt ggt gaa ggt tgt gcc agt		1767	
	Cys Gln Leu Pro Cys Pro Lys Gly Gln Phe Gly Glu Gly Cys Ala Ser			
	270	275	280	

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Val Cys Asp Cys Asp His Ser Asp Gly Cys Asp Pro Val His Gly His	
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Cys Arg Cys Gln Ala Gly Trp Met Gly Thr Arg Cys His Leu Pro Cys	
300 305 310	
cca gag ggc ttt tgg gga gcc aac tgc agc aat gcc tgt acc tgc aag	1911
Pro Glu Gly Phe Trp Gly Ala Asn Cys Ser Asn Ala Cys Thr Cys Lys	
315 320 325	
aat ggt ggc act tgt gta cct gag aac ggc aac tgt gtg tgc gca cca	1959
Asn Gly Gly Thr Cys Val Pro Glu Asn Gly Asn Cys Val Cys Ala Pro	
330 335 340 345	
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Gly Phe Arg Gly Pro Ser Cys Gln Arg Pro Cys Pro Pro Gly Arg Tyr	
350 355 360	
ggc aaa cgc tgt gtg ccc tgc aag tgc aac aac cat tct tcc tgc cac	2055
Gly Lys Arg Cys Val Pro Cys Lys Cys Asn Asn His Ser Ser Cys His	
365 370 375	
ccg tcg gat ggg acc tgc tcc tgc ctg gca ggc tgg aca ggc cct gac	2103
Pro Ser Asp Gly Thr Cys Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp	
380 385 390	
tgc tct gaa tca tgt ccc cca ggc cac tgg gga ctc aaa tgc tcc caa	2151
Cys Ser Glu Ser Cys Pro Pro Gly His Trp Gly Leu Lys Cys Ser Gln	
395 400 405	
ccc tgc cag tgt cat cat ggt gcc acc tgc cac ccc cag gat ggg agc	2199
Pro Cys Gln Cys His His Gly Ala Thr Cys His Pro Gln Asp Gly Ser	
410 415 420 425	
tgt gtc tgc atc cca ggc tgg act gga ccc aac tgc tcg gaa ggc tgc	2247
Cys Val Cys Ile Pro Gly Trp Thr Gly Pro Asn Cys Ser Glu Gly Cys	
430 435 440	
cca tca aga atg ttt ggt gtc aac tgc tcc cag cta tgt cag tgt gat	2295
Pro Ser Arg Met Phe Gly Val Asn Cys Ser Gln Leu Cys Gln Cys Asp	
445 450 455	
cct gga gag atg tgc cac cca gag act ggg gct tgc gtc tgt ccc cca	2343
Pro Gly Glu Met Cys His Pro Glu Thr Gly Ala Cys Val Cys Pro Pro	
460 465 470	
gga cac agt ggt ggc cac tgc aaa gtg ggc agc cag gag tcc ttc acc	2391
Gly His Ser Gly Ala His Cys Lys Val Gly Ser Gln Glu Ser Phe Thr	
475 480 485	
ata atg ccc acc tct cct gtg atc cat aac tca ctg ggt gcc gtg att	2439
Ile Met Pro Thr Ser Pro Val Ile His Asn Ser Leu Gly Ala Val Ile	
490 495 500 505	
ggc att gca gtg ctg ggg acc ctt gtg gtg gcc ctg gta gca ctg ttt	2487
Gly Ile Ala Val Leu Gly Thr Leu Val Val Ala Leu Val Ala Leu Phe	

510	515	520	
att ggc tac cga cac tgg caa aag ggc aag gaa cat gag cac ttg gca			2535
Ile Gly Tyr Arg His Trp Gln Lys Gly Lys Glu His Glu His Leu Ala			
525	530	535	
gtg gct tac agc act ggg cga ctg gat ggc tcc gat tac gtc atg cca			2583
Val Ala Tyr Ser Thr Gly Arg Leu Asp Gly Ser Asp Tyr Val Met Pro			
540	545	550	
gat gtc tct ccg agc tac agt cac tac tat tcc aac cct agc tac cac			2631
Asp Val Ser Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His			
555	560	565	
aca ctg tct cag tgt tct cct aac cct cca ccc cct aac aag att cca			2679
Thr Leu Ser Gln Cys Ser Pro Asn Pro Pro Pro Pro Asn Lys Ile Pro			
570	575	580	585
ggc agt cag ctg ttt gtc agc tcc cag gca tct gag cgg cca aac aga			2727
Gly Ser Gln Leu Phe Val Ser Ser Gln Ala Ser Glu Arg Pro Asn Arg			
590	595	600	
aac cat ggg cga gat aac cac gcc aca ctg ccc gct gac tgg aag cac			2775
Asn His Gly Arg Asp Asn His Ala Thr Leu Pro Ala Asp Trp Lys His			
605	610	615	
cga cgg gag tcc cat gac aga gct ttc ctc agg cac cag cca cct gga			2823
Arg Arg Glu Ser His Asp Arg Ala Phe Leu Arg His Gln Pro Pro Gly			
620	625	630	
ccg aag gta tagctgtagc tatggccaca ggaatggccc gggggccattc			2872
Pro Lys Val			
635			
tgatcataaag gtcccatctc tgaagaagga ctaggggcaa gcgttatgtc cctgagcagt			2932
gagaaccctc atgcgacctc ccgagacctg ccgggcctgc ctggggaacc ccgagaaagc			2992
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 Phe Thr Gly Gln Cys His Cys Ala Pro Gly Tyr Ile Gly Asp Arg Cys

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<220>
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 <222> (217)...(684)

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tggacaccac ctcagcccac tgagcaggag tcacagcacg aagaccaagc gcaaagcgac      180
ccctgcccctc catcctgact gctcctccta agagag atg gca ccg gcc aga gca      234
                               Met Ala Pro Ala Arg Ala
                               1           5

gga ttc tgc ccc ctt ctg ctg ctt ctg ctg ctg ggg ctg tgg gtg gca      282
Gly Phe Cys Pro Leu Leu Leu Leu Leu Leu Leu Gly Leu Trp Val Ala
                               10           15           20

gag atc cca gtc agt gcc aag ccc aag ggc atg acc tca tca cag tgg      330
Glu Ile Pro Val Ser Ala Lys Pro Lys Gly Met Thr Ser Ser Gln Trp
                               25           30           35

ttt aaa att cag cac atg cag ccc agc cct caa gca tgc aac tca gcc      378
Phe Lys Ile Gln His Met Gln Pro Ser Pro Gln Ala Cys Asn Ser Ala
                               40           45           50

atg aaa aac att aac aag cac aca aaa cgg tgc aaa gac ctc aac acc      426
Met Lys Asn Ile Asn Lys His Thr Lys Arg Cys Lys Asp Leu Asn Thr
                               55           60           65           70

ttc ctg cac gag cct ttc tcc agt gtg gcc gcc acc tgc cag acc ccc      474
Phe Leu His Glu Pro Phe Ser Ser Val Ala Ala Thr Cys Gln Thr Pro
                               75           80           85

aaa ata gcc tgc aag aat ggc gat aaa aac tgc cac cag agc cac ggg      522
Lys Ile Ala Cys Lys Asn Gly Asp Lys Asn Cys His Gln Ser His Gly
                               90           95           100

ccc gtg tcc ctg acc atg tgt aag ctc acc tca ggg aag tat ccg aac      570
Pro Val Ser Leu Thr Met Cys Lys Leu Thr Ser Gly Lys Tyr Pro Asn
                               105          110          115

tgc agg tac aaa gag aag cga cag aac aag tct tac gta gtg gcc tgt      618
Cys Arg Tyr Lys Glu Lys Arg Gln Asn Lys Ser Tyr Val Val Ala Cys
                               120          125          130

aag cct ccc cag aaa aag gac tct cag caa ttc cac ctg gtt cct gta      666
Lys Pro Pro Gln Lys Lys Asp Ser Gln Gln Phe His Leu Val Pro Val
                               135          140          145          150

cac ttg gac aga gtc ctt taggtttcca gactggcttg ctctttggct      714
His Leu Asp Arg Val Leu
                               155

gaccttcaat tccctctoca ggactccgca ccactcccct acaoccagag cattctcttc      774
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<210> 23
 <211> 156
 <212> PRT
 <213> Homo sapiens

<400> 23

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Leu	Gly	Leu	Trp	Val	Ala	Glu	Ile	Pro	Val	Ser	Ala	Lys	Pro	Lys	Gly
			20					25					30		
Met	Thr	Ser	Ser	Gln	Trp	Phe	Lys	Ile	Gln	His	Met	Gln	Pro	Ser	Pro
			35				40					45			
Gln	Ala	Cys	Asn	Ser	Ala	Met	Lys	Asn	Ile	Asn	Lys	His	Thr	Lys	Arg
			50				55				60				
Cys	Lys	Asp	Leu	Asn	Thr	Phe	Leu	His	Glu	Pro	Phe	Ser	Ser	Val	Ala
65				70					75					80	
Ala	Thr	Cys	Gln	Thr	Pro	Lys	Ile	Ala	Cys	Lys	Asn	Gly	Asp	Lys	Asn
				85					90					95	
Cys	His	Gln	Ser	His	Gly	Pro	Val	Ser	Leu	Thr	Met	Cys	Lys	Leu	Thr
			100					105				110			
Ser	Gly	Lys	Tyr	Pro	Asn	Cys	Arg	Tyr	Lys	Glu	Lys	Arg	Gln	Asn	Lys
			115				120					125			
Ser	Tyr	Val	Val	Ala	Cys	Lys	Pro	Pro	Gln	Lys	Lys	Asp	Ser	Gln	Gln
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Phe	His	Leu	Val	Pro	Val	His	Leu	Asp	Arg	Val	Leu				
145					150					155					

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 <211> 468
 <212> DNA
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<400> 24

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attcagcaca	tgcaagccag	ccctcaagca	tgcaactcag	ccatgaaaaa	cattaacaag	180
cacacaaaac	ggtgcaaaaga	cctcaacacc	ttcctgcacg	agcctttctc	cagtgtggcc	240
gccacctgcc	agacccccaa	aatagcctgc	aagaatggcg	ataaaaaactg	ccaccagagc	300
cacggggccc	tgccctgac	catgtgtaag	ctcacctcag	ggaagtatcc	gaactgcagg	360
tacaaaagaga	agcgacagaa	caagtcttac	gtagtggcct	gtaagcctcc	ccagaaaaag	420
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 <211> 1788
 <212> DNA

<213> Homo sapiens

<220>

<221> CDS

<222> (62)...(976)

<400> 25

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Met Pro Leu Leu Thr Leu Tyr Leu Leu Leu Phe Trp Leu Ser Gly Tyr
  1             5             10             15

tcc att gcc act caa atc acc ggt cca aca aca gtg aat ggc ttg gag      157
Ser Ile Ala Thr Gln Ile Thr Gly Pro Thr Thr Val Asn Gly Leu Glu
          20             25             30

cgg ggc tcc ttg acc gtg cag tgt gtt tac aga tca ggc tgg gag acc      205
Arg Gly Ser Leu Thr Val Gln Cys Val Tyr Arg Ser Gly Trp Glu Thr
          35             40             45

tac ttg aag tgg tgg tgt cga gga gct att tgg cgt gac tgc aag atc      253
Tyr Leu Lys Trp Trp Cys Arg Gly Ala Ile Trp Arg Asp Cys Lys Ile
          50             55             60

ctt gtt aaa acc agt ggg tca gag cag gag gtg aag agg gac cgg gtg      301
Leu Val Lys Thr Ser Gly Ser Glu Gln Glu Val Lys Arg Asp Arg Val
          65             70             75             80

tcc atc aag gac aat cag aaa aac cgc acg ttc act gtg acc atg gag      349
Ser Ile Lys Asp Asn Gln Lys Asn Arg Thr Phe Thr Val Thr Met Glu
          85             90             95

gat ctc atg aaa act gat gct gac act tac tgg tgt gga att gag aaa      397
Asp Leu Met Lys Thr Asp Ala Asp Thr Tyr Trp Cys Gly Ile Glu Lys
          100             105             110

act gga aat gac ctt ggg gtc aca gtt caa gtg acc att gac cca gcg      445
Thr Gly Asn Asp Leu Gly Val Thr Val Gln Val Thr Ile Asp Pro Ala
          115             120             125

tcg act cct gcc ccc acc acg cct act tcc act acg ttt aca gca cca      493
Ser Thr Pro Ala Pro Thr Thr Pro Thr Ser Thr Thr Phe Thr Ala Pro
          130             135             140

gtc acc caa gaa gaa act agc agc tcc cca act ctg acc ggc cac cac      541
Val Thr Gln Glu Glu Thr Ser Ser Ser Pro Thr Leu Thr Gly His His
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Leu Asp Asn Arg His Lys Leu Leu Lys Leu Ser Val Leu Leu Pro Leu
          165             170             175

atc ttc acc ata ttg ctg ctg ctt ttg gtg gcc gcc tca ctc ttg gct      637
Ile Phe Thr Ile Leu Leu Leu Leu Leu Val Ala Ala Ser Leu Leu Ala
          180             185             190

tgg agg atg atg aag tac cag cag aaa gca gcc ggg atg tcc cca gag      685
Trp Arg Met Met Lys Tyr Gln Gln Lys Ala Ala Gly Met Ser Pro Glu
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195	200	205	
cag gta ctg cag ccc ctg gag ggc gac ctc tgc tat gca gac ctg acc			733
Gln Val Leu Gln Pro Leu Glu Gly Asp Leu Cys Tyr Ala Asp Leu Thr			
210	215	220	
ctg cag ctg gcc gga acc tcc ccg cga aag gct acc acg aag ctt tcc			781
Leu Gln Leu Ala Gly Thr Ser Pro Arg Lys Ala Thr Thr Lys Leu Ser			
225	230	235	240
tct gcc cag gtt gac cag gtg gaa gtg gaa tat gtc acc atg gct tcc			829
Ser Ala Gln Val Asp Gln Val Glu Val Glu Tyr Val Thr Met Ala Ser			
	245	250	255
ttg ccg aag gag gac att tcc tat gca tct ctg acc ttg ggt gct gag			877
Leu Pro Lys Glu Asp Ile Ser Tyr Ala Ser Leu Thr Leu Gly Ala Glu			
	260	265	270
gat cag gaa ccg acc tac tgc aac atg ggc cac ctc agt agc cac ctc			925
Asp Gln Glu Pro Thr Tyr Cys Asn Met Gly His Leu Ser Ser His Leu			
	275	280	285
ccc ggc agg ggc cct gag gag ccc acg gaa tac agc acc atc agc agg			973
Pro Gly Arg Gly Pro Glu Glu Pro Thr Glu Tyr Ser Thr Ile Ser Arg			
	290	295	300
cct tagcctgcac tccaggctcc ttcttgacc ccaggctgtg agcacactcc			1026
Pro			
305			
tgctcatcg accgtctgcc cctgtctccc ctcatcagga ccaaccggg gactggtgcc			1086
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aatggattcc cctgcctgga gcctccaaaa gaaaccagcc ctgcccacgc cttgacttga			1686
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 Arg Gly Ser Leu Thr Val Gln Cys Val Tyr Arg Ser Gly Trp Glu Thr
 35 40 45
 Tyr Leu Lys Trp Trp Cys Arg Gly Ala Ile Trp Arg Asp Cys Lys Ile
 50 55 60

Leu Val Lys Thr Ser Gly Ser Glu Gln Glu Val Lys Arg Asp Arg Val
 65 70 75 80
 Ser Ile Lys Asp Asn Gln Lys Asn Arg Thr Phe Thr Val Thr Met Glu
 85 90 95
 Asp Leu Met Lys Thr Asp Ala Asp Thr Tyr Trp Cys Gly Ile Glu Lys
 100 105 110
 Thr Gly Asn Asp Leu Gly Val Thr Val Gln Val Thr Ile Asp Pro Ala
 115 120 125
 Ser Thr Pro Ala Pro Thr Thr Pro Thr Ser Thr Thr Phe Thr Ala Pro
 130 135 140
 Val Thr Gln Glu Glu Thr Ser Ser Ser Pro Thr Leu Thr Gly His His
 145 150 155 160
 Leu Asp Asn Arg His Lys Leu Leu Lys Leu Ser Val Leu Leu Pro Leu
 165 170 175
 Ile Phe Thr Ile Leu Leu Leu Leu Val Ala Ala Ser Leu Leu Ala
 180 185 190
 Trp Arg Met Met Lys Tyr Gln Gln Lys Ala Ala Gly Met Ser Pro Glu
 195 200 205
 Gln Val Leu Gln Pro Leu Glu Gly Asp Leu Cys Tyr Ala Asp Leu Thr
 210 215 220
 Leu Gln Leu Ala Gly Thr Ser Pro Arg Lys Ala Thr Thr Lys Leu Ser
 225 230 235 240
 Ser Ala Gln Val Asp Gln Val Glu Val Glu Tyr Val Thr Met Ala Ser
 245 250 255
 Leu Pro Lys Glu Asp Ile Ser Tyr Ala Ser Leu Thr Leu Gly Ala Glu
 260 265 270
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 Pro Gly Arg Gly Pro Glu Glu Pro Thr Glu Tyr Ser Thr Ile Ser Arg
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 Pro
 305

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<212> DNA
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Ala Leu Phe Leu Cys Leu Cys Leu Leu Thr Leu Gln Asn Ala Thr Thr
              10              15              20

gag aca tgg gaa gaa ctc ctg agc tac atg gag aat atg cag gtg tcc      152
Glu Thr Trp Glu Glu Leu Leu Ser Tyr Met Glu Asn Met Gln Val Ser
              25              30              35

agg ggc cgg agc tca gtt ttt tcc tct cgt caa ctc cac cag ctg gag      200
Arg Gly Arg Ser Ser Val Phe Ser Ser Arg Gln Leu His Gln Leu Glu
              40              45              50

cag atg cta ctg aac acc agc ttc cca ggc tac aac ctg acc ttg cag      248
Gln Met Leu Leu Asn Thr Ser Phe Pro Gly Tyr Asn Leu Thr Leu Gln
              55              60              65

aca ccc acc atc cag tct ctg gcc ttc aag ctg agc tgt gac ttc tct      296
Thr Pro Thr Ile Gln Ser Leu Ala Phe Lys Leu Ser Cys Asp Phe Ser
              70              75              80              85

ggc ctc tcg ctg acc agt gcc act ctg aag cgg gtg ccc cag gca gga      344
Gly Leu Ser Leu Thr Ser Ala Thr Leu Lys Arg Val Pro Gln Ala Gly
              90              95              100

ggt cag cat gcc cgg ggt cag cac gcc atg cag ttc ccc gcc gag ctg      392
Gly Gln His Ala Arg Gly Gln His Ala Met Gln Phe Pro Ala Glu Leu
              105              110              115

acc cgg gac gcc tgc aag acc cgc ccc agg gag ctg cgg ctc atc tgt      440
Thr Arg Asp Ala Cys Lys Thr Arg Pro Arg Glu Leu Arg Leu Ile Cys
              120              125              130

atc tac ttc tcc aac acc cac ttt ttc aag gat gaa aac aac tca tct      488
Ile Tyr Phe Ser Asn Thr His Phe Phe Lys Asp Glu Asn Asn Ser Ser
              135              140              145

ctg ctg aat aac tac gtc ctg ggg gcc cag ctg agt cat ggg cac gtg      536
Leu Leu Asn Asn Tyr Val Leu Gly Ala Gln Leu Ser His Gly His Val
              150              155              160              165

aac aac ctc agg gat cct gtg aac atc agc ttc tgg cac aac caa agc      584
Asn Asn Leu Arg Asp Pro Val Asn Ile Ser Phe Trp His Asn Gln Ser
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ctg gaa ggc tac acc ctg acc tgt gtc ttc tgg aag gag gga gcc agg      632
Leu Glu Gly Tyr Thr Leu Thr Cys Val Phe Trp Lys Glu Gly Ala Arg

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Lys Gln Pro Trp Gly Gly Trp Ser Pro Glu Gly Cys Arg Thr Glu Gln				
	200	205	210	
ccc tcc cac tct cag gtg ctc tgc cgc tgc aac cac ctc acc tac ttt				728
Pro Ser His Ser Gln Val Leu Cys Arg Cys Asn His Leu Thr Tyr Phe				
	215	220	225	
gct gtt ctc atg caa ctc tcc cca gcc ctg gtc cct gca gag ttg ctg				776
Ala Val Leu Met Gln Leu Ser Pro Ala Leu Val Pro Ala Glu Leu Leu				
	230	235	240	245
gca cct ctt acg tac atc tcc ctc gtg ggc tgc agc atc tcc atc gtg				824
Ala Pro Leu Thr Tyr Ile Ser Leu Val Gly Cys Ser Ile Ser Ile Val				
		250	255	260
gcc tog ctg atc aca gtc ctg ctg cac ttc cat ttc agg aag cag agt				872
Ala Ser Leu Ile Thr Val Leu Leu His Phe His Phe Arg Lys Gln Ser				
	265	270	275	
gac tcc tta aca cgc atc cac atg aac ctg cat gcc tcc gtg ctg ctc				920
Asp Ser Leu Thr Arg Ile His Met Asn Leu His Ala Ser Val Leu Leu				
	280	285	290	
ctg aac atc gcc ttc ctg ctg agc ccc gca ttc gca atg tct cct gtg				968
Leu Asn Ile Ala Phe Leu Leu Ser Pro Ala Phe Ala Met Ser Pro Val				
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ccc ggg tca gca tgc acg gct ctg gcc gct gcc ctg cac tac gcg ctg				1016
Pro Gly Ser Ala Cys Thr Ala Leu Ala Ala Ala Leu His Tyr Ala Leu				
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ctc agc tgc ctc acc tgg atg gcc atc gag ggc ttc aac ctc tac ctc				1064
Leu Ser Cys Leu Thr Trp Met Ala Ile Glu Gly Phe Asn Leu Tyr Leu				
		330	335	340
ctc ctc ggg cgt gtc tac aac atc tac atc cgc aga tat gtg ttc aag				1112
Leu Leu Gly Arg Val Tyr Asn Ile Tyr Ile Arg Arg Tyr Val Phe Lys				
	345	350	355	
ctt ggt gtg cta ggc tgg ggg gcc cca gcc ctc ctg gtg ctg ctt tcc				1160
Leu Gly Val Leu Gly Trp Gly Ala Pro Ala Leu Leu Val Leu Leu Ser				
	360	365	370	
ctc tct gtc aag agc tgc gta tac gga ccc tgc aca atc ccc gtc ttc				1208
Leu Ser Val Lys Ser Ser Val Tyr Gly Pro Cys Thr Ile Pro Val Phe				
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gac agc tgg gag aat ggc aca ggc ttc cag aac atg tcc ata tgc tgg				1256
Asp Ser Trp Glu Asn Gly Thr Gly Phe Gln Asn Met Ser Ile Cys Trp				
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gtg cgg agc ccc gtg gtg cac agt gtc ctg gtc atg ggc tac ggc ggc				1304
Val Arg Ser Pro Val Val His Ser Val Leu Val Met Gly Tyr Gly Gly				
	410	415	420	

ctc acg tcc ctc ttc aac ctg gtg gtg ctg gcc tgg gcg ctg tgg acc	1352
Leu Thr Ser Leu Phe Asn Leu Val Val Leu Ala Trp Ala Leu Trp Thr	
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Leu Arg Arg Leu Arg Glu Arg Ala Asp Ala Pro Ser Val Arg Ala Cys	
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His Asp Thr Val Thr Val Leu Gly Leu Thr Val Leu Leu Gly Thr Thr	
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Trp Ala Leu Ala Phe Phe Ser Phe Gly Val Phe Leu Leu Pro Gln Leu	
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Phe Leu Phe Thr Ile Leu Asn Ser Leu Tyr Gly Phe Phe Leu Phe Leu	
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Trp Phe Cys Ser Gln Arg Cys Arg Ser Glu Ala Glu Ala Lys Ala Gln	
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Ile Glu Ala Phe Ser Ser Ser Gln Thr Thr Gln	
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<210> 29

<211> 528

<212> PRT
 <213> Homo sapiens

<400> 29
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 Asn Met Gln Val Ser Arg Gly Arg Ser Ser Val Phe Ser Ser Arg Gln
 35 40 45
 Leu His Gln Leu Glu Gln Met Leu Leu Asn Thr Ser Phe Pro Gly Tyr
 50 55 60
 Asn Leu Thr Leu Gln Thr Pro Thr Ile Gln Ser Leu Ala Phe Lys Leu
 65 70 75 80
 Ser Cys Asp Phe Ser Gly Leu Ser Leu Thr Ser Ala Thr Leu Lys Arg
 85 90 95
 Val Pro Gln Ala Gly Gly Gln His Ala Arg Gly Gln His Ala Met Gln
 100 105 110
 Phe Pro Ala Glu Leu Thr Arg Asp Ala Cys Lys Thr Arg Pro Arg Glu
 115 120 125
 Leu Arg Leu Ile Cys Ile Tyr Phe Ser Asn Thr His Phe Phe Lys Asp
 130 135 140
 Glu Asn Asn Ser Ser Leu Leu Asn Asn Tyr Val Leu Gly Ala Gln Leu
 145 150 155 160
 Ser His Gly His Val Asn Asn Leu Arg Asp Pro Val Asn Ile Ser Phe
 165 170 175
 Trp His Asn Gln Ser Leu Glu Gly Tyr Thr Leu Thr Cys Val Phe Trp
 180 185 190
 Lys Glu Gly Ala Arg Lys Gln Pro Trp Gly Gly Trp Ser Pro Glu Gly
 195 200 205
 Cys Arg Thr Glu Gln Pro Ser His Ser Gln Val Leu Cys Arg Cys Asn
 210 215 220
 His Leu Thr Tyr Phe Ala Val Leu Met Gln Leu Ser Pro Ala Leu Val
 225 230 235 240
 Pro Ala Glu Leu Leu Ala Pro Leu Thr Tyr Ile Ser Leu Val Gly Cys
 245 250 255
 Ser Ile Ser Ile Val Ala Ser Leu Ile Thr Val Leu Leu His Phe His
 260 265 270
 Phe Arg Lys Gln Ser Asp Ser Leu Thr Arg Ile His Met Asn Leu His
 275 280 285
 Ala Ser Val Leu Leu Leu Asn Ile Ala Phe Leu Leu Ser Pro Ala Phe
 290 295 300
 Ala Met Ser Pro Val Pro Gly Ser Ala Cys Thr Ala Leu Ala Ala Ala
 305 310 315 320
 Leu His Tyr Ala Leu Leu Ser Cys Leu Thr Trp Met Ala Ile Glu Gly
 325 330 335
 Phe Asn Leu Tyr Leu Leu Leu Gly Arg Val Tyr Asn Ile Tyr Ile Arg
 340 345 350
 Arg Tyr Val Phe Lys Leu Gly Val Leu Gly Trp Gly Ala Pro Ala Leu
 355 360 365
 Leu Val Leu Leu Ser Leu Ser Val Lys Ser Ser Val Tyr Gly Pro Cys
 370 375 380
 Thr Ile Pro Val Phe Asp Ser Trp Glu Asn Gly Thr Gly Phe Gln Asn
 385 390 395 400
 Met Ser Ile Cys Trp Val Arg Ser Pro Val Val His Ser Val Leu Val
 405 410 415
 Met Gly Tyr Gly Gly Leu Thr Ser Leu Phe Asn Leu Val Val Leu Ala
 420 425 430

Trp Ala Leu Trp Thr Leu Arg Arg Leu Arg Glu Arg Ala Asp Ala Pro
 435 440 445
 Ser Val Arg Ala Cys His Asp Thr Val Thr Val Leu Gly Leu Thr Val
 450 455 460
 Leu Leu Gly Thr Thr Trp Ala Leu Ala Phe Phe Ser Phe Gly Val Phe
 465 470 475 480
 Leu Leu Pro Gln Leu Phe Leu Phe Thr Ile Leu Asn Ser Leu Tyr Gly
 485 490 495
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 <211> 1584
 <212> DNA
 <213> Homo sapiens

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<210> 31
 <211> 63
 <212> PRT
 <213> Homo sapiens

<400> 31
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 Lys Asp Leu Phe Leu Cys His Pro Glu Phe Lys Ser Gly Glu Tyr Trp
 20 25 30
 Ile Asp Pro Asn Gln Gly Cys Ile Lys Asp Ala Ile Lys Val Phe Cys

<210> 32
<211> 25
<212> PRT
<213> Homo sapiens

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<210> 33
<211> 33
<212> PRT
<213> Homo sapiens

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<210> 34
<211> 65
<212> PRT
<213> Homo sapiens
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<210> 35
<211> 26
<212> PRT
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<212> PRT
 <213> Homo sapiens

<400> 36
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 1 5 10 15
 Ala Thr Phe Leu Phe His Thr Gln Glu Pro Asn Gln Leu Pro Val Ile
 20 25 30

<210> 37
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 37
 Gly Glu Ser Val Thr Leu Thr Cys Ser Val Ser Gly Phe Gly Pro Pro
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 Pro Val Thr Trp Leu Arg Asn Gly Lys Leu Ser Leu Thr Ile Ser
 20 25 30

<210> 38
 <211> 57
 <212> PRT
 <213> Homo sapiens

<400> 38
 Gly Arg Thr Val Arg Leu Gln Cys Pro Val Glu Gly Asp Pro Pro Pro
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 Thr Met Trp Thr Lys Asp Gly Arg Thr Ile His Ser Gly Trp Ser Arg
 20 25 30
 Phe Arg Val Leu Pro Gln Gly Leu Lys Val Lys Gln Val Glu Arg Glu
 35 40 45
 Asp Ala Gly Val Tyr Val Cys Lys Ala
 50 55

<210> 39
 <211> 59
 <212> PRT
 <213> Homo sapiens

<400> 39
 Gly Ser Ser Val Arg Leu Lys Cys Val Ala Ser Gly His Pro Arg Pro
 1 5 10 15
 Asp Ile Thr Trp Met Lys Asp Asp Gln Ala Leu Thr Arg Pro Glu Ala
 20 25 30
 Ala Glu Pro Arg Lys Lys Lys Trp Thr Leu Ser Leu Lys Asn Leu Arg
 35 40 45
 Pro Glu Asp Ser Gly Lys Tyr Thr Cys Arg Val
 50 55

<210> 40
 <211> 79
 <212> PRT
 <213> Homo sapiens

<400> 40
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 1 5 10 15

Val Ile Gln Trp Leu Lys Arg Val Glu Tyr Gly Ala Glu Gly Arg His
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Asn Ser Thr Ile Asp Val Gly Gly Gln Lys Phe Val Val Leu Pro Thr
35 40 45
Gly Asp Val Trp Ser Arg Pro Asp Gly Ser Tyr Asn Lys Leu Leu Ile
50 55 60
Thr Arg Ala Arg Gln Asp Asp Ala Gly Met Tyr Ile Cys Leu Gly
65 70 75

<210> 41
<211> 78
<212> PRT
<213> Homo sapiens

<400> 41
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Tyr Leu Lys Trp Trp Cys Arg Gly Ala Ile Trp Arg Asp Cys Lys Ile
20 25 30
Leu Val Lys Thr Ser Gly Ser Glu Gln Glu Val Lys Arg Asp Arg Val
35 40 45
Ser Ile Lys Asp Asn Gln Lys Asn Arg Thr Phe Thr Val Thr Met Glu
50 55 60
Asp Leu Met Lys Thr Asp Ala Asp Thr Tyr Trp Cys Gly Ile
65 70 75

<210> 42
<211> 10
<212> PRT
<213> Homo sapiens

<400> 42
Val Phe Val Leu Gly Thr Leu Gly Ile Phe
1 5 10

<210> 43
<211> 10
<212> PRT
<213> Homo sapiens

<400> 43
Val Phe Ile Leu Gly Thr Leu Leu Leu Trp
1 5 10

<210> 44
<211> 116
<212> PRT
<213> Homo sapiens

<400> 44
Cys Gly Gly Thr Leu Asp Leu Thr Glu Ser Ser Gly Ser Ile Ser Ser
1 5 10 15
Pro Asn Tyr Pro Asn Arg Ser Asp Tyr Pro Pro Asn Lys Glu Cys Val
20 25 30
Trp Arg Ile Arg Ala Pro Pro Gly Tyr Arg Val Val Glu Leu Thr Phe
35 40 45
Gln Asp Phe Asp Leu Glu Asp His Asp Gly Ala Pro Cys Arg Tyr Asp
50 55 60

Tyr Val Glu Ile Arg Asp Gly Asp Pro Ser Ser Pro Leu Leu Gly Arg
 65 70 75 80
 Phe Cys Gly Ser Gly Lys Pro Glu Asp Ile Arg Ser Thr Ser Asn Arg
 85 90 95
 Met Leu Ile Lys Phe Val Ser Asp Ala Ser Val Ser Lys Arg Gly Phe
 100 105 110
 Lys Ala Thr Tyr
 115

<210> 45
 <211> 97
 <212> PRT
 <213> Homo sapiens

<400> 45
 Gly Ser Val Leu Leu Ala Gln Glu Leu Pro Gln Gln Leu Thr Ser Pro
 1 5 10 15
 Gly Tyr Pro Glu Pro Tyr Gly Lys Gly Gln Glu Ser Ser Thr Asp Ile
 20 25 30
 Lys Ala Pro Glu Gly Phe Ala Val Arg Leu Val Phe Gln Asp Phe Asp
 35 40 45
 Leu Glu Pro Ser Gln Asp Cys Ala Gly Asp Ser Val Thr Val Ser Trp
 50 55 60
 Gly Trp Gly Gly Ser Arg Gln Asp Cys Gly Gln Gly Asp Ser Arg Gly
 65 70 75 80
 Cys Gly Lys Trp Arg Cys Pro Glu Ser Pro Ile Trp Arg Arg Asp Glu
 85 90 95
 Phe

<210> 46
 <211> 45
 <212> PRT
 <213> Homo sapiens

<400> 46
 Cys Ala Pro Asn Asn Pro Cys Ser Asn Gly Gly Thr Cys Val Asn Thr
 1 5 10 15
 Pro Gly Gly Ser Ser Asp Asn Phe Gly Gly Tyr Thr Cys Glu Cys Pro
 20 25 30
 Pro Gly Asp Tyr Tyr Leu Ser Tyr Thr Gly Lys Arg Cys
 35 40 45

<210> 47
 <211> 67
 <212> PRT
 <213> Homo sapiens

<400> 47
 Trp Ser Thr Asp Lys His Ile Gly Gly Arg Thr Ser Leu Gly Phe Asn
 1 5 10 15
 Leu Glu Tyr Arg Ile Arg Val Thr Cys Asp Glu Asn Tyr Tyr Gly Glu
 20 25 30
 Gly Cys Asn Lys Phe Cys Arg Pro Arg Asp Ala Phe Gly His Tyr
 35 40 45
 Thr Cys Asp Glu Asn Gly Asn Lys Leu Cys Leu Glu Gly Trp Lys Gly
 50 55 60
 Glu Tyr Cys

65

<210> 48
 <211> 59
 <212> PRT
 <213> Homo sapiens

<400> 48
 Cys Asp Cys Asn Pro His Gly Ser Leu Ser Asp Asp Thr Cys Asp Ser
 1 5 10 15
 Asp Asp Glu Leu Phe Gly Glu Glu Thr Gly Gln Cys Leu Lys Cys Lys
 20 25 30
 Pro Asn Val Thr Gly Arg Arg Cys Asp Arg Cys Lys Pro Gly Tyr Tyr
 35 40 45
 Gly Leu Pro Ser Gly Asp Pro Gln Gln Gly Cys
 50 55

<210> 49
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 49
 Cys Val Pro Leu Cys Ala Gln Glu Cys Val His Gly Arg Cys Val Ala
 1 5 10 15
 Pro Asn Gln Cys Gln Cys Val Pro Gly Trp Arg Gly Asp Asp Cys
 20 25 30

<210> 50
 <211> 30
 <212> PRT
 <213> Homo sapiens

<400> 50
 Cys Gln Phe Arg Cys Gln Cys His Gly Ala Pro Cys Asp Pro Gln Thr
 1 5 10 15
 Gly Ala Cys Phe Cys Pro Ala Glu Arg Thr Gly Pro Ser Cys
 20 25 30

<210> 51
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 51
 Cys Pro Ser Thr His Pro Cys Gln Asn Gly Gly Val Phe Gln Thr Pro
 1 5 10 15
 Gln Gly Ser Cys Ser Cys Pro Pro Gly Trp Met Gly Thr Ile Cys
 20 25 30

<210> 52
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 52
 Cys Ser Gln Glu Cys Arg Cys His Asn Gly Gly Leu Cys Asp Arg Phe
 1 5 10 15

Thr Gly Gln Cys Arg Cys Ala Pro Gly Tyr Thr Gly Asp Arg Cys
 20 25 30

<210> 53
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 53
 Cys Ala Glu Thr Cys Asp Cys Ala Pro Asp Ala Arg Cys Phe Pro Ala
 1 5 10 15
 Asn Gly Ala Cys Leu Cys Glu His Gly Phe Thr Gly Asp Arg Cys
 20 25 30

<210> 54
 <211> 27
 <212> PRT
 <213> Homo sapiens

<400> 54
 Cys Asp Arg Glu His Ser Leu Ser Cys His Pro Met Asn Gly Glu Cys
 1 5 10 15
 Ser Cys Leu Pro Gly Trp Ala Gly Leu His Cys
 20 25

<210> 55
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 55
 Cys Gln Glu His Cys Leu Cys Leu His Gly Gly Val Cys Gln Ala Thr
 1 5 10 15
 Ser Gly Leu Cys Gln Cys Ala Pro Gly Tyr Thr Gly Pro His Cys
 20 25 30

<210> 56
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 56
 Cys Ser Ala Arg Cys Ser Cys Glu Asn Ala Ile Ala Cys Ser Pro Ile
 1 5 10 15
 Asp Gly Glu Cys Val Cys Lys Glu Gly Trp Gln Arg Gly Asn Cys
 20 25 30

<210> 57
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 57
 Cys Asn Ala Ser Cys Gln Cys Ala His Glu Ala Val Cys Ser Pro Gln
 1 5 10 15
 Thr Gly Ala Cys Thr Cys Thr Pro Gly Trp His Gly Ala His Cys
 20 25 30

<210> 58
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 58
 Cys Ala Ser Arg Cys Asp Cys Asp His Ser Asp Gly Cys Asp Pro Val
 1 5 10 15
 His Gly Arg Cys Gln Cys Gln Ala Gly Trp Met Gly Ala Arg Cys
 20 25 30

<210> 59
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 59
 Cys Ser Asn Thr Cys Thr Cys Lys Asn Gly Gly Thr Cys Leu Pro Glu
 1 5 10 15
 Asn Gly Asn Cys Val Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys
 20 25 30

<210> 60
 <211> 30
 <212> PRT
 <213> Homo sapiens

<400> 60
 Cys Val Pro Cys Lys Cys Ala Asn His Ser Phe Cys His Pro Ser Asn
 1 5 10 15
 Gly Thr Cys Tyr Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys
 20 25 30

<210> 61
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 61
 Cys Ala Gln Thr Cys Gln Cys His His Gly Gly Thr Cys His Pro Gln
 1 5 10 15
 Asp Gly Ser Cys Ile Cys Pro Leu Gly Trp Thr Gly His His Cys
 20 25 30

<210> 62
 <211> 31
 <212> PRT
 <213> Homo sapiens

<400> 62
 Cys Ser Gln Pro Cys Gln Cys Gly Pro Gly Glu Lys Cys His Pro Glu
 1 5 10 15
 Thr Gly Ala Cys Val Cys Pro Pro Gly His Ser Gly Ala Pro Cys
 20 25 30

<210> 63
 <211> 37
 <212> PRT

<213> Homo sapiens

<400> 63

Gln Thr Gly Ala Cys Thr Cys Thr Pro Gly Trp His Gly Ala His Cys
1 5 10 15
Gln Leu Pro Cys Pro Lys Gly Gln Phe Gly Glu Gly Cys Ala Ser Arg
20 25 30
Cys Asp Cys Asp His
35

<210> 64

<211> 31

<212> PRT

<213> Mus musculus

<400> 64

Cys Ser Asn Thr Cys Thr Cys Lys Asn Gly Gly Thr Cys Val Ser Glu
1 5 10 15
Asn Gly Asn Cys Val Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys
20 25 30

<210> 65

<211> 31

<212> PRT

<213> Mus musculus

<400> 65

Cys Val Gln Cys Lys Cys Asn Asn Asn His Ser Ser Cys His Pro Ser
1 5 10 15
Asp Gly Thr Cys Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys
20 25 30

<210> 66

<211> 31

<212> PRT

<213> Mus musculus

<400> 66

Cys Ser Gln Leu Cys Gln Cys His His Gly Gly Thr Cys His Pro Gln
1 5 10 15
Asp Gly Ser Cys Ile Cys Thr Pro Gly Trp Thr Gly Pro Asn Cys
20 25 30

<210> 67

<211> 31

<212> PRT

<213> Mus musculus

<400> 67

Cys Ser Gln Leu Cys Gln Cys Asp Leu Gly Glu Met Cys His Pro Glu
1 5 10 15
Thr Gly Ala Cys Val Cys Pro Pro Gly His Ser Gly Ala Asp Cys
20 25 30

<210> 68

<211> 35

<212> PRT

<213> Mus musculus

<400> 68
 His Ala Ser Gly Asp Pro Val His Gly Gln Cys Arg Cys Gln Ala Gly
 1 5 10 15
 Trp Met Gly Thr Arg Cys His Leu Pro Cys Pro Glu Gly Phe Trp Gly
 20 25 30
 Ala Asn Cys
 35

<210> 69
 <211> 40
 <212> PRT
 <213> Mus musculus

<400> 69
 Cys Thr Cys Lys Asn Gly Gly Thr Cys Val Ser Glu Asn Gly Asn Cys
 1 5 10 15
 Val Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys Gln Arg Pro Cys Pro
 20 25 30
 Pro Gly Arg Tyr Gly Lys Arg Cys
 35 40

<210> 70
 <211> 35
 <212> PRT
 <213> Mus musculus

<400> 70
 Cys Lys Cys Asn Asn His Ser Ser Cys His Pro Ser Asp Gly Thr
 1 5 10 15
 Cys Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys Ser Glu Ala Cys
 20 25 30
 Pro Pro Gly
 35

<210> 71
 <211> 34
 <212> PRT
 <213> Mus musculus

<400> 71
 Cys Gln Cys His His Gly Gly Thr Cys His Pro Gln Asp Gly Ser Cys
 1 5 10 15
 Ile Cys Thr Pro Gly Trp Thr Gly Pro Asn Cys Leu Glu Gly Cys Pro
 20 25 30
 Pro Arg

<210> 72
 <211> 58
 <212> PRT
 <213> Mus musculus

<400> 72
 His Gly Gln Cys Arg Cys Gln Ala Gly Trp Met Gly Thr Arg Cys His
 1 5 10 15
 Leu Pro Cys Pro Glu Gly Phe Trp Gly Ala Asn Cys Ser Asn Thr Cys
 20 25 30
 Thr Cys Lys Asn Gly Gly Thr Cys Val Ser Glu Asn Gly Asn Cys Val

35 40 45
 Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys
 50 55

<210> 73
 <211> 28
 <212> PRT
 <213> Rattus sp.

<400> 73
 Glu Cys Arg Cys His Asn Gly Gly Leu Cys Asp Arg Phe Thr Gly Gln
 1 5 10 15
 Cys His Cys Ala Pro Gly Tyr Ile Gly Asp Arg Cys
 20 25

<210> 74
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 74
 Cys Ala Glu Thr Cys Asp Cys Ala Pro Gly Ala Arg Cys Phe Pro Ala
 1 5 10 15
 Asn Gly Ala Cys Leu Cys Glu His Gly Phe Thr Gly Asp Arg Cys
 20 25 30

<210> 75
 <211> 33
 <212> PRT
 <213> Rattus sp.

<400> 75
 Cys Gln Asp Pro Cys Thr Cys Asp Pro Glu His Ser Leu Ser Cys His
 1 5 10 15
 Pro Met His Gly Glu Cys Ser Cys Gln Pro Gly Trp Ala Gly Leu His
 20 25 30
 Cys

<210> 76
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 76
 Cys Gln Glu His Cys Leu Cys Leu His Gly Gly Val Cys Leu Ala Asp
 1 5 10 15
 Ser Gly Leu Cys Arg Cys Ala Pro Gly Tyr Thr Gly Pro His Cys
 20 25 30

<210> 77
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 77
 Cys Ser Ser His Cys Ser Cys Glu Asn Ala Ile Ala Cys Ser Pro Val
 1 5 10 15

Asp Gly Thr Cys Ile Cys Lys Glu Gly Trp Gln Arg Gly Asn Cys
 20 25 30

<210> 78
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 78
 Cys Asn Ala Ser Cys Gln Cys Ala His Glu Gly Val Cys Ser Pro Gln
 1 5 10 15
 Thr Gly Ala Cys Thr Cys Thr Pro Gly Trp Arg Gly Val His Cys
 20 25 30

<210> 79
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 79
 Cys Ala Ser Val Cys Asp Cys Asp His Ser Asp Gly Cys Asp Pro Val
 1 5 10 15
 His Gly His Cys Arg Cys Gln Ala Gly Trp Met Gly Thr Arg Cys
 20 25 30

<210> 80
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 80
 Cys Ser Asn Ala Cys Thr Cys Lys Asn Gly Gly Thr Cys Val Pro Glu
 1 5 10 15
 Asn Gly Asn Cys Val Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys
 20 25 30

<210> 81
 <211> 30
 <212> PRT
 <213> Rattus sp.

<400> 81
 Cys Val Pro Cys Lys Cys Asn Asn His Ser Ser Cys His Pro Ser Asp
 1 5 10 15
 Gly Thr Cys Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys
 20 25 30

<210> 82
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 82
 Cys Ser Gln Pro Cys Gln Cys His His Gly Ala Thr Cys His Pro Gln
 1 5 10 15
 Asp Gly Ser Cys Val Cys Ile Pro Gly Trp Thr Gly Pro Asn Cys
 20 25 30

<210> 83
 <211> 31
 <212> PRT
 <213> Rattus sp.

<400> 83
 Cys Ser Gln Leu Cys Gln Cys Asp Pro Gly Glu Met Cys His Pro Glu
 1 5 10 15
 Thr Gly Ala Cys Val Cys Pro Pro Gly His Ser Gly Ala His Cys
 20 25 30

<210> 84
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 84
 Cys Arg Cys His Asn Gly Gly Leu Cys Asp Arg Phe Thr Gly Gln Cys
 1 5 10 15
 His Cys Ala Pro Gly Tyr Ile Gly Asp Arg Cys Arg Glu Glu Cys Pro
 20 25 30
 Val Gly Arg Phe Gly Gln Asp Cys
 35 40

<210> 85
 <211> 39
 <212> PRT
 <213> Rattus sp.

<400> 85
 Cys Asp Cys Ala Pro Gly Ala Arg Cys Phe Pro Ala Asn Gly Ala Cys
 1 5 10 15
 Leu Cys Glu His Gly Phe Thr Gly Asp Arg Cys Thr Glu Arg Leu Cys
 20 25 30
 Pro Asp Gly Tyr Gly Leu Cys
 35

<210> 86
 <211> 42
 <212> PRT
 <213> Rattus sp.

<400> 86
 Cys Thr Cys Asp Pro Glu His Ser Leu Ser Cys His Pro Met His Gly
 1 5 10 15
 Glu Cys Ser Cys Gln Pro Gly Trp Ala Gly Leu His Cys Asn Glu Ser
 20 25 30
 Cys Pro Gln Asp Thr His Gly Ala Gly Cys
 35 40

<210> 87
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 87
 Cys Leu Cys Leu His Gly Gly Val Cys Leu Ala Asp Ser Gly Leu Cys
 1 5 10 15

Arg Cys Ala Pro Gly Tyr Thr Gly Pro His Cys Ala Asn Leu Cys Pro
 20 25 30
 Pro Asn Thr Tyr Gly Ile Asn Cys
 35 40

<210> 88
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 88
 Cys Ser Cys Glu Asn Ala Ile Ala Cys Ser Pro Val Asp Gly Thr Cys
 1 5 10 15
 Ile Cys Lys Glu Gly Trp Gln Arg Gly Asn Cys Ser Val Pro Cys Pro
 20 25 30
 Pro Gly Thr Trp Gly Phe Ser Cys
 35 40

<210> 89
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 89
 Cys Gln Cys Ala His Glu Gly Val Cys Ser Pro Gln Thr Gly Ala Cys
 1 5 10 15
 Thr Cys Thr Pro Gly Trp Arg Gly Val His Cys Gln Leu Pro Cys Pro
 20 25 30
 Lys Gly Gln Phe Gly Glu Gly Cys
 35 40

<210> 90
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 90
 Cys Asp Cys Asp His Ser Asp Gly Cys Asp Pro Val His Gly His Cys
 1 5 10 15
 Arg Cys Gln Ala Gly Trp Met Gly Thr Arg Cys His Leu Pro Cys Pro
 20 25 30
 Glu Gly Phe Trp Gly Ala Asn Cys
 35 40

<210> 91
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 91
 Cys Thr Cys Lys Asn Gly Gly Thr Cys Val Pro Glu Asn Gly Asn Cys
 1 5 10 15
 Val Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys Gln Arg Pro Cys Pro
 20 25 30
 Pro Gly Arg Tyr Gly Lys Arg Cys
 35 40

<210> 92

<211> 40
 <212> PRT
 <213> Rattus sp.

<400> 92
 Cys Lys Cys Asn Asn His Ser Ser Cys His Pro Ser Asp Gly Thr Cys
 1 5 10 15
 Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys Ser Glu Ser Cys Pro
 20 25 30
 Pro Gly His Trp Gly Leu Lys Cys
 35 40

<210> 93
 <211> 40
 <212> PRT
 <213> Rattus sp.

<400> 93
 Cys Gln Cys His His Gly Ala Thr Cys His Pro Gln Asp Gly Ser Cys
 1 5 10 15
 Val Cys Ile Pro Gly Trp Thr Gly Pro Asn Cys Ser Glu Gly Cys Pro
 20 25 30
 Ser Arg Met Phe Gly Val Asn Cys
 35 40

<210> 94
 <211> 36
 <212> PRT
 <213> Rattus sp.

<400> 94
 Cys Gln Cys Asp Pro Gly Glu Met Cys His Pro Glu Thr Gly Ala Cys
 1 5 10 15
 Val Cys Pro Pro Gly His Ser Gly Ala His Cys Lys Val Gly Ser Gln
 20 25 30
 Glu Ser Phe Thr
 35

<210> 95
 <211> 64
 <212> PRT
 <213> Rattus sp.

<400> 95
 Gly Val Cys Ser Pro Gln Thr Gly Ala Cys Thr Cys Thr Pro Gly Trp
 1 5 10 15
 Arg Gly Val His Cys Gln Leu Pro Cys Pro Lys Gly Gln Phe Gly Glu
 20 25 30
 Gly Cys Ala Ser Val Cys Asp Cys Asp His Ser Asp Gly Cys Asp Pro
 35 40 45
 Val His Gly His Cys Arg Cys Gln Ala Gly Trp Met Gly Thr Arg Cys
 50 55 60

<210> 96
 <211> 129
 <212> PRT
 <213> Homo sapiens


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<400> 96
Gln Glu Ser Arg Ala Gln Lys Phe Leu Arg Gln His Ile Asp Ser Pro
 1          5          10          15
Lys Thr Ser Ser Asn Pro Asn Tyr Cys Asn Gln Met Met Asp Lys
          20          25          30
Arg Arg Asn Met Thr Gln Gln Arg Cys Lys Pro Val Asn Thr Phe Val
          35          40          45
His Glu Ser Leu Ala Asp Val Lys Ala Val Cys Ser Gln Lys Asn Val
 50          55          60
Thr Cys Lys Asn Gly Gln Ser Lys Ser Ser Phe Gln Ile Thr Asp Cys
65          70          75          80
Arg Leu Thr Gly Gly Ser Gln Lys Tyr Pro Asn Cys Arg Tyr Arg Thr
          85          90          95
Ser Ala Ser Thr Lys His Ile Ile Val Ala Cys Glu Gly Arg Asp Arg
          100          105          110
Asp Asp Pro Tyr Tyr Asn Pro Tyr Val Pro Val His Phe Asp Ala Ser
          115          120          125
Val

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<210> 97
<211> 125
<212> PRT
<213> Homo sapiens

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<400> 97
Gly Met Thr Ser Ser Gln Trp Phe Lys Ile Gln His Met Gln Pro Ser
 1          5          10          15
Pro Gln Ala Cys Asn Ser Ala Met Lys Asn Ile Asn Lys His Thr Lys
          20          25          30
Arg Cys Lys Asp Leu Asn Thr Phe Leu His Glu Pro Phe Ser Ser Val
          35          40          45
Ala Ala Thr Cys Gln Thr Pro Lys Ile Ala Cys Lys Asn Gly Asp Lys
 50          55          60
Asn Cys His Gln Ser His Gly Pro Val Ser Leu Thr Met Cys Lys Leu
65          70          75          80
Thr Ser Gly Lys Tyr Pro Asn Cys Arg Tyr Lys Glu Lys Arg Gln Asn
          85          90          95
Lys Ser Tyr Val Val Ala Cys Lys Pro Pro Gln Lys Lys Asp Ser Gln
          100          105          110
Gln Phe His Leu Val Pro Val His Leu Asp Arg Val Leu
          115          120          125

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<210> 98
<211> 411
<212> PRT
<213> Homo sapiens

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<400> 98
Cys Asn Arg Thr Trp Asp Gly Ile Thr Cys Trp Pro Asp Thr Pro Pro
 1          5          10          15
Gly Glu Leu Val Val Val Pro Cys Pro Lys Tyr Phe Tyr Gly Phe Ser
          20          25          30
Ser Asp Gln Thr Asp Thr Thr Gly Asn Val Ser Arg Asn Cys Thr Glu
          35          40          45
Asp Gly Ser Trp Ser Glu Pro Pro Ser Asn Arg Thr Trp Arg Asn
 50          55          60
Tyr Ser Ala Cys Gly Glu Asp Asp Pro Glu Glu Glu Ser Glu Lys Lys

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65					70					75				80	
Lys	Lys	Tyr	Tyr	Leu	Val	Leu	Lys	Ile	Ile	Tyr	Thr	Val	Gly	Tyr	Ser
				85					90					95	
Leu	Ser	Leu	Ala	Ala	Leu	Leu	Val	Ala	Val	Val	Ile	Leu	Leu	Leu	Phe
			100					105					110		
Arg	Lys	Leu	His	Thr	Leu	Trp	Pro	Asp	Asn	Ala	Asp	Gly	Ala	Leu	Glu
		115					120					125			
Val	Gly	Ala	Pro	Trp	Gly	Ala	Pro	Phe	Gln	Val	Arg	Arg	Ser	Ile	Arg
	130				135					140					
Cys	Thr	Arg	Asn	Tyr	Ile	His	Met	Asn	Leu	Phe	Leu	Ser	Phe	Ile	Leu
145					150					155					160
Arg	Ala	Ala	Ser	Val	Phe	Ile	Lys	Asp	Ala	Val	Leu	Lys	Ser	Glu	Val
			165					170						175	
Ser	Ser	Asp	Glu	Pro	Glu	Arg	Leu	Ser	Ser	Arg	Cys	Ser	Leu	Ser	Thr
			180					185					190		
Gly	Gln	Val	Val	Val	Gly	Cys	Lys	Leu	Leu	Val	Val	Phe	Gln	Phe	Gln
	195						200					205			
Tyr	Cys	Val	Met	Thr	Asn	Phe	Phe	Trp	Leu	Leu	Val	Glu	Gly	Leu	Tyr
	210				215						220				
Leu	His	Thr	Leu	Leu	Val	Val	Thr	Phe	Phe	Ser	Glu	Arg	Lys	Tyr	Leu
225					230					235					240
Trp	Trp	Tyr	Leu	Leu	Ile	Gly	Trp	Gly	Val	Pro	Leu	Val	Phe	Val	Thr
			245						250					255	
Val	Trp	Ala	Ile	Val	Arg	Leu	Leu	Phe	Glu	Asp	Thr	Gly	Cys	Trp	Asp
			260					265					270		
Ser	Asn	Gly	Leu	Ala	Met	Phe	Pro	Glu	Ala	Lys	Met	Cys	Ile	Trp	Met
		275					280					285			
Ser	Asp	Asn	Ser	His	Leu	Trp	Trp	Ile	Ile	Lys	Gly	Pro	Ile	Leu	Leu
	290				295					300					
Ser	Ile	Leu	Val	Asn	Phe	Phe	Leu	Phe	Ile	Asn	Ile	Ile	Arg	Ile	Leu
305				310						315					320
Val	Thr	Lys	Leu	Arg	Ala	Ala	Gln	Thr	Gly	Glu	Thr	Asp	Gln	Arg	Gln
			325						330				335		
Tyr	Ser	Gln	Tyr	Arg	Lys	Leu	Ala	Lys	Ser	Thr	Leu	Leu	Leu	Ile	Pro
			340					345					350		
Leu	Phe	Gly	Ile	His	Tyr	Val	Val	Phe	Ala	Phe	Arg	Pro	Ser	Asn	Asp
		355					360					365			
Ala	Arg	Gly	Val	Leu	Arg	Lys	Ile	Lys	Leu	Tyr	Phe	Glu	Leu	Ser	Leu
	370					375					380				
Gly	Ser	Phe	Gln	Gly	Phe	Phe	Val	Ala	Val	Leu	Tyr	Cys	Phe	Leu	Asn
385					390					395					400
Gly	Glu	Val	Gln	Ala	Glu	Ile	Arg	Arg	Arg	Trp					
			405						410						

<210> 99
 <211> 328
 <212> PRT
 <213> Homo sapiens

<400> 99
 Leu Thr Cys Val Phe Trp Lys Glu Gly Ala Arg Lys Gln Pro Trp Gly
 1 5 10 15
 Gly Trp Ser Pro Glu Gly Cys Arg Thr Glu Gln Pro Ser His Ser Gln
 20 25 30
 Val Leu Cys Arg Cys Asn His Leu Thr Tyr Phe Ala Val Leu Met Gln
 35 40 45
 Leu Ser Pro Ala Leu Val Pro Ala Glu Leu Leu Ala Pro Leu Thr Tyr
 50 55 60

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Ile Ser Leu Val Gly Cys Ser Ile Ser Ile Val Ala Ser Leu Ile Thr
65      70      75      80
Val Leu Leu His Phe Arg Lys Gln Ser Asp Ser Leu Thr Arg Ile His
      85      90      95
Met Asn Leu His Ala Ser Val Leu Leu Leu Asn Ile Ala Phe Leu Leu
      100      105      110
Ser Pro Ala Phe Ala Met Ser Pro Val Pro Gly Ser Ala Cys Thr Ala
      115      120      125
Leu Ala Ala Ala Leu His Tyr Ala Leu Leu Ser Cys Leu Thr Trp Met
      130      135      140
Ala Ile Glu Gly Phe Asn Leu Tyr Leu Leu Leu Gly Arg Val Tyr Asn
      145      150      155      160
Ile Tyr Ile Arg Arg Tyr Val Phe Lys Leu Gly Val Leu Gly Trp Gly
      165      170      175
Ala Pro Ala Leu Leu Val Leu Leu Ser Leu Ser Val Lys Ser Ser Val
      180      185      190
Tyr Gly Pro Cys Thr Ile Pro Val Phe Asp Ser Trp Glu Asn Gly Thr
      195      200      205
Gly Phe Gln Asn Met Ser Ile Cys Trp Val Arg Ser Pro Val Val His
      210      215      220
Ser Val Leu Val Met Gly Tyr Gly Gly Leu Thr Ser Leu Phe Asn Leu
      225      230      235      240
Val Val Leu Ala Trp Ala Leu Trp Thr Leu Arg Arg Leu Arg Glu Arg
      245      250      255
Ala Asp Ala Pro Ser Val Arg Ala Cys His Asp Thr Val Thr Val Leu
      260      265      270
Gly Leu Thr Val Leu Leu Gly Thr Thr Trp Ala Leu Ala Phe Phe Ser
      275      280      285
Phe Gly Val Phe Leu Leu Pro Gln Leu Phe Leu Phe Thr Ile Leu Asn
      290      295      300
Ser Leu Tyr Gly Phe Phe Leu Phe Leu Trp Phe Cys Ser Gln Arg Cys
      305      310      315      320
Arg Ser Glu Ala Glu Ala Lys Ala
      325

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<210> 100
<211> 150
<212> PRT
<213> Pan troglodytes

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<400> 100
Met Val Leu Cys Phe Pro Leu Leu Leu Leu Leu Val Leu Trp Gly
1      5      10      15
Pro Val Cys Pro Leu His Ala Trp Pro Lys Arg Leu Thr Lys Ala His
      20      25      30
Trp Phe Glu Ile Gln His Ile Gln Pro Ser Pro Leu Gln Cys Asn Arg
      35      40      45
Ala Met Ser Gly Ile Asn Asn Tyr Ala Gln His Cys Lys His Gln Asn
      50      55      60
Thr Phe Leu His Asp Ser Phe Gln Asn Val Ala Ala Val Cys Asp Leu
      65      70      75      80
Leu Ser Ile Val Cys Lys Asn Arg Arg His Asn Cys His Gln Ser Ser
      85      90      95
Lys Pro Val Asn Met Thr Asp Cys Arg Leu Thr Ser Gly Lys Tyr Pro
      100      105      110
Gln Cys Arg Tyr Ser Ala Ala Ala Gln Tyr Lys Phe Phe Ile Val Ala
      115      120      125
Cys Asp Pro Pro Gln Lys Ser Asp Pro Pro Tyr Lys Leu Val Pro Val

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      130              135              140
His Leu Asp Ser Ile Leu
145              150

      <210> 101
      <211> 24
      <212> PRT
      <213> Homo sapiens

      <400> 101
Met Thr Pro Ser Pro Leu Leu Leu Leu Leu Leu Pro Pro Leu Leu Leu
 1          5          10          15
Gly Ala Phe Pro Pro Ala Ala Ala

      <210> 102
      <211> 480
      <212> PRT
      <213> Homo sapiens

      <400> 102
Ala Arg Gly Pro Pro Lys Met Ala Asp Lys Val Val Pro Arg Gln Val
 1          5          10          15
Ala Arg Leu Gly Arg Thr Val Arg Leu Gln Cys Pro Val Glu Gly Asp
 20          25          30
Pro Pro Pro Leu Thr Met Trp Thr Lys Asp Gly Arg Thr Ile His Ser
 35          40          45
Gly Trp Ser Arg Phe Arg Val Leu Pro Gln Gly Leu Lys Val Lys Gln
 50          55          60
Val Glu Arg Glu Asp Ala Gly Val Tyr Val Cys Lys Ala Thr Asn Gly
 65          70          75          80
Phe Gly Ser Leu Ser Val Asn Tyr Thr Leu Val Val Leu Asp Asp Ile
 85          90          95
Ser Pro Gly Lys Glu Ser Leu Gly Pro Asp Ser Ser Ser Gly Gly Gln
 100         105         110
Glu Asp Pro Ala Ser Gln Gln Trp Ala Arg Pro Arg Phe Thr Gln Pro
 115         120         125
Ser Lys Met Arg Arg Arg Val Ile Ala Arg Pro Val Gly Ser Ser Val
 130         135         140
Arg Leu Lys Cys Val Ala Ser Gly His Pro Arg Pro Asp Ile Thr Trp
 145         150         155         160
Met Lys Asp Asp Gln Ala Leu Thr Arg Pro Glu Ala Ala Glu Pro Arg
 165         170         175
Lys Lys Lys Trp Thr Leu Ser Leu Lys Asn Leu Arg Pro Glu Asp Ser
 180         185         190
Gly Lys Tyr Thr Cys Arg Val Ser Asn Arg Ala Gly Ala Ile Asn Ala
 195         200         205
Thr Tyr Lys Val Asp Val Ile Gln Arg Thr Arg Ser Lys Pro Val Leu
 210         215         220
Thr Gly Thr His Pro Val Asn Thr Thr Val Asp Phe Gly Gly Thr Thr
 225         230         235         240
Ser Phe Gln Cys Lys Val Arg Ser Asp Val Lys Pro Val Ile Gln Trp
 245         250         255
Leu Lys Arg Val Glu Tyr Gly Ala Glu Gly Arg His Asn Ser Thr Ile
 260         265         270
Asp Val Gly Gly Gln Lys Phe Val Val Leu Pro Thr Gly Asp Val Trp
 275         280         285
Ser Arg Pro Asp Gly Ser Tyr Leu Asn Lys Leu Leu Ile Thr Arg Ala

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290		295		300
Arg	Gln	Asp	Asp	Ala
305		310		315
Gly	Tyr	Ser	Phe	Arg
		325		330
Pro	Pro	Gly	Pro	Pro
		340		345
Trp	Pro	Val	Val	Ile
		355		360
Thr	Leu	Leu	Leu	Trp
370		375		380
Ala	Pro	Ala	Pro	Pro
385		390		395
Asp	Arg	Ser	Gly	Asp
		405		410
Gly	Pro	Gly	Val	Gly
		420		425
Gln	His	Leu	Gly	Pro
		435		440
Lys	Leu	Tyr	Thr	Asp
450		455		460
His	Ser	His	Val	Glu
465		470		475

<210> 103
 <211> 350
 <212> PRT
 <213> Homo sapiens

<400> 103
Ala Arg Gly Pro Pro Lys Met Ala Asp Lys Val Val Pro Arg Gln Val
1 5 10 15
Ala Arg Leu Gly Arg Thr Val Arg Leu Gln Cys Pro Val Glu Gly Asp
20 25 30
Pro Pro Pro Leu Thr Met Trp Thr Lys Asp Gly Arg Thr Ile His Ser
35 40 45
Gly Trp Ser Arg Phe Arg Val Leu Pro Gln Gly Leu Lys Val Lys Gln
50 55 60
Val Glu Arg Glu Asp Ala Gly Val Tyr Val Cys Lys Ala Thr Asn Gly
65 70 75 80
Phe Gly Ser Leu Ser Val Asn Tyr Thr Leu Val Val Leu Asp Asp Ile
85 90 95
Ser Pro Gly Lys Glu Ser Leu Gly Pro Asp Ser Ser Ser Gly Gly Gln
100 105 110
Glu Asp Pro Ala Ser Gln Gln Trp Ala Arg Pro Arg Phe Thr Gln Pro
115 120 125
Ser Lys Met Arg Arg Arg Val Ile Ala Arg Pro Val Gly Ser Ser Val
130 135 140
Arg Leu Lys Cys Val Ala Ser Gly His Pro Arg Pro Asp Ile Thr Trp
145 150 155 160
Met Lys Asp Asp Gln Ala Leu Thr Arg Pro Glu Ala Ala Glu Pro Arg
165 170 175
Lys Lys Lys Trp Thr Leu Ser Leu Lys Asn Leu Arg Pro Glu Asp Ser
180 185 190
Gly Lys Tyr Thr Cys Arg Val Ser Asn Arg Ala Gly Ala Ile Asn Ala
195 200 205
Thr Tyr Lys Val Asp Val Ile Gln Arg Thr Arg Ser Lys Pro Val Leu
210 215 220

Thr Gly Thr His Pro Val Asn Thr Thr Val Asp Phe Gly Gly Thr Thr
 225 230 235 240
 Ser Phe Gln Cys Lys Val Arg Ser Asp Val Lys Pro Val Ile Gln Trp
 245 250 255
 Leu Lys Arg Val Glu Tyr Gly Ala Glu Gly Arg His Asn Ser Thr Ile
 260 265 270
 Asp Val Gly Gly Gln Lys Phe Val Val Leu Pro Thr Gly Asp Val Trp
 275 280 285
 Ser Arg Pro Asp Gly Ser Tyr Leu Asn Lys Leu Leu Ile Thr Arg Ala
 290 295 300
 Arg Gln Asp Asp Ala Gly Met Tyr Ile Cys Leu Gly Ala Asn Thr Met
 305 310 315 320
 Gly Tyr Ser Phe Arg Ser Ala Phe Leu Thr Val Leu Pro Asp Pro Lys
 325 330 335
 Pro Pro Gly Pro Pro Val Ala Ser Ser Ser Ser Ala Thr Ser
 340 345 350

<210> 104
 <211> 24
 <212> PRT
 <213> Homo sapiens

<400> 104
 Leu Pro Trp Pro Val Val Ile Gly Ile Pro Ala Gly Ala Val Phe Ile
 1 5 10 15
 Leu Gly Thr Leu Leu Leu Trp Leu
 20

<210> 105
 <211> 106
 <212> PRT
 <213> Homo sapiens

<400> 105
 Cys Gln Ala Gln Lys Lys Pro Cys Thr Pro Ala Pro Ala Pro Pro Leu
 1 5 10 15
 Pro Gly His Arg Pro Pro Gly Thr Ala Arg Asp Arg Ser Gly Asp Lys
 20 25 30
 Asp Leu Pro Ser Leu Ala Ala Leu Ser Ala Gly Pro Gly Val Gly Leu
 35 40 45
 Cys Glu Glu His Gly Ser Pro Ala Ala Pro Gln His Leu Leu Gly Pro
 50 55 60
 Gly Pro Val Ala Gly Pro Lys Leu Tyr Pro Lys Leu Tyr Thr Asp Ile
 65 70 75 80
 His Thr His Thr His Thr His Ser His Thr His Ser His Val Glu Gly
 85 90 95
 Lys Val His Gln His Ile His Tyr Gln Cys
 100 105

<210> 106
 <211> 208
 <212> PRT
 <213> Mus musculus

<400> 106
 Arg Val Arg Pro Thr Gly Asp Val Trp Ser Arg Pro Asp Gly Ser Tyr
 1 5 10 15
 Leu Asn Lys Leu Leu Ile Ser Arg Ala Arg Gln Asp Asp Ala Gly Met

```

      20      25      30
Tyr Ile Cys Leu Gly Ala Asn Thr Met Gly Tyr Ser Phe Arg Ser Ala
      35      40      45
Phe Leu Thr Val Leu Pro Asp Pro Lys Pro Pro Gly Pro Pro Met Ala
      50      55      60
Ser Ser Ser Ser Ser Thr Ser Leu Pro Trp Pro Val Val Ile Gly Ile
      65      70      75      80
Pro Ala Gly Ala Val Phe Ile Leu Gly Thr Val Leu Leu Trp Leu Cys
      85      90      95
Gln Thr Lys Lys Lys Pro Cys Ala Pro Ala Ser Thr Leu Pro Val Pro
      100      105      110
Gly His Arg Pro Pro Gly Thr Ser Arg Glu Arg Ser Gly Asp Lys Asp
      115      120      125
Leu Pro Ser Leu Ala Val Gly Ile Cys Glu Glu His Gly Ser Ala Met
      130      135      140
Ala Pro Gln His Ile Leu Ala Ser Gly Ser Thr Ala Gly Pro Lys Leu
      145      150      155      160
Tyr Pro Lys Leu Tyr Thr Asp Val His Thr His Thr His Thr Thr
      165      170      175
Cys Thr His Thr Leu Ser Cys Trp Arg Ala Arg Phe Ile Asn Thr Ser
      180      185      190      195
Met Ser Thr Ile Ser Ala Lys Tyr Ser Glu Ser Pro Ser Thr Val Ser
      195      200      205

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```

<210> 107
<211> 73
<212> PRT
<213> Mus musculus

```

```

      <400> 107
Arg Val Arg Pro Thr Gly Asp Val Trp Ser Arg Pro Asp Gly Ser Tyr
      1      5      10      15
Leu Asn Lys Leu Leu Ile Ser Arg Ala Arg Gln Asp Asp Ala Gly Met
      20      25      30
Tyr Ile Cys Leu Gly Ala Asn Thr Met Gly Tyr Ser Phe Arg Ser Ala
      35      40      45
Phe Leu Thr Val Leu Pro Asp Pro Lys Pro Pro Gly Pro Pro Met Ala
      50      55      60
Ser Ser Ser Ser Ser Thr Ser Leu Pro
      65      70

```

```

<210> 108
<211> 23
<212> PRT
<213> Mus musculus

```

```

      <400> 108
Trp Pro Val Val Ile Gly Ile Pro Ala Gly Ala Val Phe Ile Leu Gly
      1      5      10      15
Thr Val Leu Leu Trp Leu Cys
      20

```

```

<210> 109
<211> 112
<212> PRT
<213> Mus musculus

```

```

<400> 109

```

Gln Thr Lys Lys Lys Pro Cys Ala Pro Ala Ser Thr Leu Pro Val Pro
 1 5 10 15
 Gly His Arg Pro Pro Gly Thr Ser Arg Glu Arg Ser Gly Asp Lys Asp
 20 25 30
 Leu Pro Ser Leu Ala Val Gly Ile Cys Glu Glu His Gly Ser Ala Met
 35 40 45
 Ala Pro Gln His Ile Leu Ala Ser Gly Ser Thr Ala Gly Pro Lys Leu
 50 55 60
 Tyr Pro Lys Leu Tyr Thr Asp Val His Thr His Thr His Thr His Thr
 65 70 75 80
 Cys Thr His Thr Leu Ser Cys Trp Arg Ala Arg Phe Ile Asn Thr Ser
 85 90 95
 Met Ser Thr Ile Ser Ala Lys Tyr Ser Glu Ser Pro Ser Thr Val Ser
 100 105 110

<210> 110
 <211> 35
 <212> PRT
 <213> Homo sapiens

<400> 110
 Met Pro Gly Pro Arg Val Trp Gly Lys Tyr Leu Trp Arg Ser Pro His
 1 5 10 15
 Ser Lys Gly Cys Pro Gly Ala Met Trp Trp Leu Leu Leu Trp Gly Val
 20 25 30
 Leu Gln Ala
 35

<210> 111
 <211> 103
 <212> PRT
 <213> Homo sapiens

<400> 111
 Cys Pro Thr Arg Gly Ser Val Leu Leu Ala Gln Glu Leu Pro Gln Gln
 1 5 10 15
 Leu Thr Ser Pro Gly Tyr Pro Glu Pro Tyr Gly Lys Gly Gln Glu Ser
 20 25 30
 Ser Thr Asp Ile Lys Ala Pro Glu Gly Phe Ala Val Arg Leu Val Phe
 35 40 45
 Gln Asp Phe Asp Leu Glu Pro Ser Gln Asp Cys Ala Gly Asp Ser Val
 50 55 60
 Thr Val Ser Trp Gly Trp Gly Gly Ser Arg Gln Asp Cys Gly Gln Gly
 65 70 75 80
 Asp Ser Arg Gly Cys Gly Lys Trp Arg Cys Pro Glu Ser Pro Ile Trp
 85 90 95
 Arg Arg Asp Glu Phe Ser Met
 100

<210> 112
 <211> 20
 <212> PRT
 <213> Homo sapiens

<400> 112
 Met Ser Pro Pro Leu Cys Pro Leu Leu Leu Leu Ala Val Gly Leu Arg
 1 5 10 15
 Leu Ala Gly Thr

20

<210> 113
 <211> 1030
 <212> PRT
 <213> Homo sapiens

<400> 113
 Leu Asn Pro Ser Asp Pro Asn Thr Cys Ser Phe Trp Glu Ser Phe Thr
 1 5 10 15
 Thr Thr Thr Lys Glu Ser His Ser Arg Pro Phe Ser Leu Leu Pro Ser
 20 25 30
 Glu Pro Cys Glu Arg Pro Trp Glu Gly Pro His Thr Cys Pro Ser Pro
 35 40 45
 Gln Thr Gln Arg Lys Leu Leu Ala Ser Arg Asp Ser Phe Cys Met Val
 50 55 60
 Cys Val Gly Ala Gly Val Gln Trp Arg Asp Arg Ser Ala Leu Gln Pro
 65 70 75 80
 Gln Thr Gly Asn Ala Leu Ser Met Arg Pro Gln Pro Arg Val Leu Ser
 85 90 95
 Gly Ala Pro Ser Leu Ala Ser Pro Gly His Thr Val Val Val Lys Thr
 100 105 110
 Asp His Arg Gln Arg Leu Gln Cys Cys His Gly Phe Tyr Glu Ser Arg
 115 120 125
 Gly Phe Cys Val Pro Leu Cys Ala Gln Glu Cys Val His Gly Arg Cys
 130 135 140
 Val Ala Pro Asn Gln Cys Gln Cys Val Pro Gly Trp Arg Gly Asp Asp
 145 150 155 160
 Cys Ser Ser Ala Pro Asn Cys Leu Gln Pro Cys Thr Pro Gly Tyr Tyr
 165 170 175
 Gly Pro Ala Cys Gln Phe Arg Cys Gln Cys His Gly Ala Pro Cys Asp
 180 185 190
 Pro Gln Thr Gly Ala Cys Phe Cys Pro Ala Glu Arg Thr Gly Pro Ser
 195 200 205
 Cys Asp Val Ser Cys Ser Gln Gly Thr Ser Gly Phe Phe Cys Pro Ser
 210 215 220
 Thr His Pro Cys Gln Asn Gly Gly Val Phe Gln Thr Pro Gln Gly Ser
 225 230 235 240
 Cys Ser Cys Pro Pro Gly Trp Met Gly Thr Ile Cys Ser Leu Pro Cys
 245 250 255
 Pro Glu Gly Phe His Gly Pro Asn Cys Ser Gln Glu Cys Arg Cys His
 260 265 270
 Asn Gly Gly Leu Cys Asp Arg Phe Thr Gly Gln Cys Arg Cys Ala Pro
 275 280 285
 Gly Tyr Thr Gly Asp Arg Cys Arg Glu Glu Cys Pro Val Gly Arg Phe
 290 295 300
 Gly Gln Asp Cys Ala Glu Thr Cys Asp Cys Ala Pro Asp Ala Arg Cys
 305 310 315 320
 Phe Pro Ala Asn Gly Ala Cys Leu Cys Glu His Gly Phe Thr Gly Asp
 325 330 335
 Arg Cys Thr Asp Arg Leu Cys Pro Asp Gly Phe Tyr Gly Leu Ser Cys
 340 345 350
 Gln Ala Pro Cys Thr Cys Asp Arg Glu His Ser Leu Ser Cys His Pro
 355 360 365
 Met Asn Gly Glu Cys Ser Cys Leu Pro Gly Trp Ala Gly Leu His Cys
 370 375 380
 Asn Glu Ser Cys Pro Gln Asp Thr His Gly Pro Gly Cys Gln Glu His
 385 390 395 400

Cys	Leu	Cys	Leu	His	Gly	Gly	Val	Cys	Gln	Ala	Thr	Ser	Gly	Leu	Cys	
				405					410					415		
Gln	Cys	Ala	Pro	Gly	Tyr	Thr	Gly	Pro	His	Cys	Ala	Ser	Leu	Cys	Pro	
			420					425					430			
Pro	Asp	Thr	Tyr	Gly	Val	Asn	Cys	Ser	Ala	Arg	Cys	Ser	Cys	Glu	Asn	
		435					440					445				
Ala	Ile	Ala	Cys	Ser	Pro	Ile	Asp	Gly	Glu	Cys	Val	Cys	Lys	Glu	Gly	
	450					455					460					
Trp	Gln	Arg	Gly	Asn	Cys	Ser	Val	Pro	Cys	Pro	Pro	Gly	Thr	Trp	Gly	
465				470						475				480		
Phe	Ser	Cys	Asn	Ala	Ser	Cys	Gln	Cys	Ala	His	Glu	Ala	Val	Cys	Ser	
			485						490					495		
Pro	Gln	Thr	Gly	Ala	Cys	Thr	Cys	Thr	Pro	Gly	Trp	His	Gly	Ala	His	
		500						505					510			
Cys	Gln	Leu	Pro	Cys	Pro	Lys	Gly	Gln	Phe	Gly	Glu	Gly	Cys	Ala	Ser	
	515					520						525				
Arg	Cys	Asp	Cys	Asp	His	Ser	Asp	Gly	Cys	Asp	Pro	Val	His	Gly	Arg	
	530				535						540					
Cys	Gln	Cys	Gln	Ala	Gly	Trp	Met	Gly	Ala	Arg	Cys	His	Leu	Ser	Cys	
545				550					555					560		
Pro	Glu	Gly	Leu	Trp	Gly	Val	Asn	Cys	Ser	Asn	Thr	Cys	Thr	Cys	Lys	
			565					570					575			
Asn	Gly	Gly	Thr	Cys	Leu	Pro	Glu	Asn	Gly	Asn	Cys	Val	Cys	Ala	Pro	
		580						585					590			
Gly	Phe	Arg	Gly	Pro	Ser	Cys	Gln	Arg	Ser	Cys	Gln	Pro	Gly	Arg	Tyr	
	595					600						605				
Gly	Lys	Arg	Cys	Val	Pro	Cys	Lys	Cys	Ala	Asn	His	Ser	Phe	Cys	His	
	610				615						620					
Pro	Ser	Asn	Gly	Thr	Cys	Tyr	Cys	Leu	Ala	Gly	Trp	Thr	Gly	Pro	Asp	
625				630						635				640		
Cys	Ser	Gln	Pro	Cys	Pro	Pro	Gly	His	Trp	Gly	Glu	Asn	Cys	Ala	Gln	
			645						650					655		
Thr	Cys	Gln	Cys	His	His	Gly	Gly	Thr	Cys	His	Pro	Gln	Asp	Gly	Ser	
		660						665					670			
Cys	Ile	Cys	Pro	Leu	Gly	Trp	Thr	Gly	His	His	Cys	Leu	Glu	Gly	Cys	
	675						680						685			
Pro	Leu	Gly	Thr	Phe	Gly	Ala	Asn	Cys	Ser	Gln	Pro	Cys	Gln	Cys	Gly	
	690					695					700					
Pro	Gly	Glu	Lys	Cys	His	Pro	Glu	Thr	Gly	Ala	Cys	Val	Cys	Pro	Pro	
705				710						715				720		
Gly	His	Ser	Gly	Ala	Pro	Cys	Arg	Ile	Gly	Ile	Gln	Glu	Pro	Phe	Thr	
			725						730					735		
Val	Met	Pro	Thr	Thr	Pro	Val	Ala	Tyr	Asn	Ser	Leu	Gly	Ala	Val	Ile	
			740					745					750			
Gly	Ile	Ala	Val	Leu	Gly	Ser	Leu	Val	Val	Ala	Leu	Val	Ala	Leu	Phe	
	755						760						765			
Ile	Gly	Tyr	Arg	His	Trp	Gln	Lys	Gly	Lys	Glu	His	His	His	Leu	Ala	
	770					775					780					
Val	Ala	Tyr	Ser	Ser	Gly	Arg	Leu	Asp	Gly	Ser	Glu	Tyr	Val	Met	Pro	
785				790						795				800		
Asp	Val	Pro	Pro	Ser	Tyr	Ser	His	Tyr	Tyr	Ser	Asn	Pro	Ser	Tyr	His	
			805						810					815		
Thr	Leu	Ser	Gln	Cys	Ser	Pro	Asn	Pro	Pro	Pro	Pro	Asn	Lys	Val	Pro	
		820						825					830			
Gly	Pro	Leu	Phe	Ala	Ser	Leu	Gln	Asn	Pro	Glu	Arg	Pro	Gly	Gly	Ala	
	835						840					845				
Gln	Gly	His	Asp	Asn	His	Thr	Thr	Leu	Pro	Ala	Asp	Trp	Lys	His	Arg	
	850					855					860					

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Arg Glu Pro Pro Pro Gly Pro Leu Asp Arg Gly Ser Ser Arg Leu Asp
865      870      875      880
Arg Ser Tyr Ser Tyr Ser Tyr Ser Asn Gly Pro Gly Pro Phe Tyr Asp
      885      890      895
Lys Gly Leu Ile Ser Glu Glu Glu Leu Gly Ala Ser Val Ala Ser Leu
      900      905      910
Ser Ser Glu Asn Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu Pro
      915      920      925
Gly Gly Pro Arg Glu Ser Ser Tyr Met Glu Met Lys Gly Pro Pro Ser
      930      935      940
Gly Ser Ala Pro Arg Gln Pro Pro Gln Phe Trp Asp Ser Gln Arg Arg
      945      950      955      960
Arg Gln Pro Gln Pro Gln Arg Asp Ser Gly Thr Tyr Glu Gln Pro Ser
      965      970      975
Pro Leu Ile His Asp Arg Asp Ser Val Gly Ser Gln Pro Pro Leu Pro
      980      985      990
Pro Gly Leu Pro Pro Gly His Tyr Asp Ser Pro Lys Asn Ser His Ile
      995      1000      1005
Pro Gly His Tyr Asp Leu Pro Pro Val Arg His Pro Pro Ser Pro Pro
      1010      1015      1020
Leu Arg Arg Gln Asp Arg
1025      1030

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<210> 114
<211> 747
<212> PRT
<213> Homo sapiens

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<400> 114
Leu Asn Pro Ser Asp Pro Asn Thr Cys Ser Phe Trp Glu Ser Phe Thr
1      5      10      15
Thr Thr Thr Lys Glu Ser His Ser Arg Pro Phe Ser Leu Leu Pro Ser
      20      25      30
Glu Pro Cys Glu Arg Pro Trp Glu Gly Pro His Thr Cys Pro Ser Pro
      35      40      45
Gln Thr Gln Arg Lys Leu Leu Ala Ser Arg Asp Ser Phe Cys Met Val
      50      55      60
Cys Val Gly Ala Gly Val Gln Trp Arg Asp Arg Ser Ala Leu Gln Pro
      65      70      75      80
Gln Thr Gly Asn Ala Leu Ser Met Arg Pro Gln Pro Arg Val Leu Ser
      85      90      95
Gly Ala Pro Ser Leu Ala Ser Pro Gly His Thr Val Val Val Lys Thr
      100      105      110
Asp His Arg Gln Arg Leu Gln Cys Cys His Gly Phe Tyr Glu Ser Arg
      115      120      125
Gly Phe Cys Val Pro Leu Cys Ala Gln Glu Cys Val His Gly Arg Cys
      130      135      140
Val Ala Pro Asn Gln Cys Gln Cys Val Pro Gly Trp Arg Gly Asp Asp
      145      150      155      160
Cys Ser Ser Ala Pro Asn Cys Leu Gln Pro Cys Thr Pro Gly Tyr Tyr
      165      170      175
Gly Pro Ala Cys Gln Phe Arg Cys Gln Cys His Gly Ala Pro Cys Asp
      180      185      190
Pro Gln Thr Gly Ala Cys Phe Cys Pro Ala Glu Arg Thr Gly Pro Ser
      195      200      205
Cys Asp Val Ser Cys Ser Gln Gly Thr Ser Gly Phe Phe Cys Pro Ser
      210      215      220
Thr His Pro Cys Gln Asn Gly Gly Val Phe Gln Thr Pro Gln Gly Ser

```

225					230					235				240
Cys	Ser	Cys	Pro	Pro	Gly	Trp	Met	Gly	Thr	Ile	Cys	Ser	Leu	Pro
				245					250					255
Pro	Glu	Gly	Phe	His	Gly	Pro	Asn	Cys	Ser	Gln	Glu	Cys	Arg	Cys
			260					265					270	
Asn	Gly	Gly	Leu	Cys	Asp	Arg	Phe	Thr	Gly	Gln	Cys	Arg	Cys	Ala
		275				280						285		Pro
Gly	Tyr	Thr	Gly	Asp	Arg	Cys	Arg	Glu	Glu	Cys	Pro	Val	Gly	Arg
	290					295					300			Phe
Gly	Gln	Asp	Cys	Ala	Glu	Thr	Cys	Asp	Cys	Ala	Pro	Asp	Ala	Arg
305					310					315				320
Phe	Pro	Ala	Asn	Gly	Ala	Cys	Leu	Cys	Glu	His	Gly	Phe	Thr	Gly
			325						330					335
Arg	Cys	Thr	Asp	Arg	Leu	Cys	Pro	Asp	Gly	Phe	Tyr	Gly	Leu	Ser
			340					345					350	Cys
Gln	Ala	Pro	Cys	Thr	Cys	Asp	Arg	Glu	His	Ser	Leu	Ser	Cys	His
	355					360						365		Pro
Met	Asn	Gly	Glu	Cys	Ser	Cys	Leu	Pro	Gly	Trp	Ala	Gly	Leu	His
	370					375				380				Cys
Asn	Glu	Ser	Cys	Pro	Gln	Asp	Thr	His	Gly	Pro	Gly	Cys	Gln	Glu
385					390					395				400
Cys	Leu	Cys	Leu	His	Gly	Gly	Val	Cys	Gln	Ala	Thr	Ser	Gly	Leu
			405						410					Cys
Gln	Cys	Ala	Pro	Gly	Tyr	Thr	Gly	Pro	His	Cys	Ala	Ser	Leu	Cys
			420					425					430	Pro
Pro	Asp	Thr	Tyr	Gly	Val	Asn	Cys	Ser	Ala	Arg	Cys	Ser	Cys	Glu
	435					440					445			Asn
Ala	Ile	Ala	Cys	Ser	Pro	Ile	Asp	Gly	Glu	Cys	Val	Cys	Lys	Glu
	450					455					460			Gly
Trp	Gln	Arg	Gly	Asn	Cys	Ser	Val	Pro	Cys	Pro	Pro	Gly	Thr	Trp
465				470					475					Gly
Phe	Ser	Cys	Asn	Ala	Ser	Cys	Gln	Cys	Ala	His	Glu	Ala	Val	Ser
			485						490				495	
Pro	Gln	Thr	Gly	Ala	Cys	Thr	Cys	Thr	Pro	Gly	Trp	His	Gly	Ala
		500						505					510	His
Cys	Gln	Leu	Pro	Cys	Pro	Lys	Gly	Gln	Phe	Gly	Glu	Gly	Cys	Ala
	515					520						525		Ser
Arg	Cys	Asp	Cys	Asp	His	Ser	Asp	Gly	Cys	Asp	Pro	Val	His	Gly
	530					535					540			Arg
Cys	Gln	Cys	Gln	Ala	Gly	Trp	Met	Gly	Ala	Arg	Cys	His	Leu	Ser
545					550				555					Cys
Pro	Glu	Gly	Leu	Trp	Gly	Val	Asn	Cys	Ser	Asn	Thr	Cys	Thr	Lys
			565						570				575	
Asn	Gly	Gly	Thr	Cys	Leu	Pro	Glu	Asn	Gly	Asn	Cys	Val	Cys	Ala
		580						585				590		Pro
Gly	Phe	Arg	Gly	Pro	Ser	Cys	Gln	Arg	Ser	Cys	Gln	Pro	Gly	Arg
	595						600					605		Tyr
Gly	Lys	Arg	Cys	Val	Pro	Cys	Lys	Cys	Ala	Asn	His	Ser	Phe	Cys
	610					615					620			His
Pro	Ser	Asn	Gly	Thr	Cys	Tyr	Cys	Leu	Ala	Gly	Trp	Thr	Gly	Pro
625					630				635					Asp
Cys	Ser	Gln	Pro	Cys	Pro	Pro	Gly	His	Trp	Gly	Glu	Asn	Cys	Ala
			645						650					Gln
Thr	Cys	Gln	Cys	His	His	Gly	Gly	Thr	Cys	His	Pro	Gln	Asp	Gly
		660						665				670		Ser
Cys	Ile	Cys	Pro	Leu	Gly	Trp	Thr	Gly	His	His	Cys	Leu	Glu	Gly
	675						680					685		Cys
Pro	Leu	Gly	Thr	Phe	Gly	Ala	Asn	Cys	Ser	Gln	Pro	Cys	Gln	Cys

690 695 700
 Pro Gly Glu Lys Cys His Pro Glu Thr Gly Ala Cys Val Cys Pro Pro
 705 710 715 720
 Gly His Ser Gly Ala Pro Cys Arg Ile Gly Ile Gln Glu Pro Phe Thr
 725 730 735
 Val Met Pro Thr Thr Pro Val Ala Tyr Asn Ser
 740 745

<210> 115
 <211> 24
 <212> PRT
 <213> Homo sapiens

<400> 115
 Leu Gly Ala Val Ile Gly Ile Ala Val Leu Gly Ser Leu Val Val Ala
 1 5 10 15
 Leu Val Ala Leu Phe Ile Gly Tyr
 20

<210> 116
 <211> 259
 <212> PRT
 <213> Homo sapiens

<400> 116
 Arg His Trp Gln Lys Gly Lys Glu His His His Leu Ala Val Ala Tyr
 1 5 10 15
 Ser Ser Gly Arg Leu Asp Gly Ser Glu Tyr Val Met Pro Asp Val Pro
 20 25 30
 Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser
 35 40 45
 Gln Cys Ser Pro Asn Pro Pro Pro Asn Lys Val Pro Gly Pro Leu
 50 55 60
 Phe Ala Ser Leu Gln Asn Pro Glu Arg Pro Gly Gly Ala Gln Gly His
 65 70 75 80
 Asp Asn His Thr Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu Pro
 85 90 95
 Pro Pro Gly Pro Leu Asp Arg Gly Ser Ser Arg Leu Asp Arg Ser Tyr
 100 105 110
 Ser Tyr Ser Tyr Ser Asn Gly Pro Gly Pro Phe Tyr Asp Lys Gly Leu
 115 120 125
 Ile Ser Glu Glu Glu Leu Gly Ala Ser Val Ala Ser Leu Ser Ser Glu
 130 135 140
 Asn Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu Pro Gly Gly Pro
 145 150 155 160
 Arg Glu Ser Ser Tyr Met Glu Met Lys Gly Pro Pro Ser Gly Ser Ala
 165 170 175
 Pro Arg Gln Pro Pro Gln Phe Trp Asp Ser Gln Arg Arg Arg Gln Pro
 180 185 190
 Gln Pro Gln Arg Asp Ser Gly Thr Tyr Glu Gln Pro Ser Pro Leu Ile
 195 200 205
 His Asp Arg Asp Ser Val Gly Ser Gln Pro Pro Leu Pro Pro Gly Leu
 210 215 220
 Pro Pro Gly His Tyr Asp Ser Pro Lys Asn Ser His Ile Pro Gly His
 225 230 235 240
 Tyr Asp Leu Pro Pro Val Arg His Pro Pro Ser Pro Pro Leu Arg Arg
 245 250 255
 Gln Asp Arg

<210> 117
 <211> 497
 <212> PRT
 <213> Mus msuculus

<400> 117
 Ser Thr His Ala Ser Gly Asp Pro Val His Gly Gln Cys Arg Cys Gln
 1 5 10 15
 Ala Gly Trp Met Gly Thr Arg Cys His Leu Pro Cys Pro Glu Gly Phe
 20 25 30
 Trp Gly Ala Asn Cys Ser Asn Thr Cys Thr Cys Lys Asn Gly Gly Thr
 35 40 45
 Cys Val Ser Glu Asn Gly Asn Cys Val Cys Ala Pro Gly Phe Arg Gly
 50 55 60
 Pro Ser Cys Gln Arg Pro Cys Pro Pro Gly Arg Tyr Gly Lys Arg Cys
 65 70 75 80
 Val Gln Cys Lys Cys Asn Asn Asn His Ser Ser Cys His Pro Ser Asp
 85 90 95
 Gly Thr Cys Ser Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys Ser Glu
 100 105 110
 Ala Cys Pro Pro Gly His Trp Gly Leu Lys Cys Ser Gln Leu Cys Gln
 115 120 125
 Cys His His Gly Gly Thr Cys His Pro Gln Asp Gly Ser Cys Ile Cys
 130 135 140
 Thr Pro Gly Trp Thr Gly Pro Asn Cys Leu Glu Gly Cys Pro Pro Arg
 145 150 155 160
 Met Phe Gly Val Asn Cys Ser Gln Leu Cys Gln Cys Asp Leu Gly Glu
 165 170 175
 Met Cys His Pro Glu Thr Gly Ala Cys Val Cys Pro Pro Gly His Ser
 180 185 190
 Gly Ala Asp Cys Lys Met Gly Ser Gln Glu Ser Phe Thr Ile Met Pro
 195 200 205
 Thr Ser Pro Val Thr His Asn Ser Leu Gly Ala Val Ile Gly Ile Ala
 210 215 220
 Val Leu Gly Thr Leu Val Ala Leu Ile Ala Leu Phe Ile Gly Tyr
 225 230 235 240
 Arg Gln Trp Gln Lys Gly Lys Glu His Glu His Leu Ala Val Ala Tyr
 245 250 255
 Ser Thr Gly Arg Leu Asp Gly Ser Asp Tyr Val Met Pro Asp Val Ser
 260 265 270
 Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser
 275 280 285
 Gln Cys Ser Pro Asn Pro Pro Pro Pro Asn Lys Val Pro Gly Ser Gln
 290 295 300
 Leu Phe Val Ser Ser Gln Ala Pro Glu Arg Pro Ser Arg Ala His Gly
 305 310 315 320
 Arg Glu Asn His Thr Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu
 325 330 335
 Pro His Asp Arg Gly Ala Ser His Leu Asp Arg Ser Tyr Ser Cys Ser
 340 345 350
 Tyr Ser His Arg Asn Gly Pro Gly Pro Phe Cys His Lys Gly Pro Ile
 355 360 365
 Ser Glu Gly Gly Leu Gly Ala Ser Val Met Ser Leu Ser Ser Glu Asn
 370 375 380
 Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu Pro Gly Glu Pro Arg
 385 390 395 400

Leu Ile Ala Leu Phe Ile Gly Tyr
20

<210> 120
<211> 257
<212> PRT
<213> Mus musculus

<400> 120
Arg Gln Trp Gln Lys Gly Lys Glu His Glu His Leu Ala Val Ala Tyr
1 5 10 15
Ser Thr Gly Arg Leu Asp Gly Ser Asp Tyr Val Met Pro Asp Val Ser
20 25 30
Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser
35 40 45
Gln Cys Ser Pro Asn Pro Pro Pro Pro Asn Lys Val Pro Gly Ser Gln
50 55 60
Leu Phe Val Ser Ser Gln Ala Pro Glu Arg Pro Ser Arg Ala His Gly
65 70 75 80
Arg Glu Asn His Thr Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu
85 90 95
Pro His Asp Arg Gly Ala Ser His Leu Asp Arg Ser Tyr Ser Cys Ser
100 105 110
Tyr Ser His Arg Asn Gly Pro Gly Pro Phe Cys His Lys Gly Pro Ile
115 120 125
Ser Glu Glu Gly Leu Gly Ala Ser Val Met Ser Leu Ser Ser Glu Asn
130 135 140
Pro Tyr Ala Thr Ile Arg Asp Leu Pro Ser Leu Pro Gly Glu Pro Arg
145 150 155 160
Glu Ser Gly Tyr Val Glu Met Lys Gly Pro Pro Ser Val Ser Pro Pro
165 170 175
Arg Gln Ser Leu His Leu Arg Asp Arg Gln Gln Arg Gln Gln Pro
180 185 190
Gln Arg Asp Ser Gly Thr Tyr Glu Gln Pro Ser Pro Leu Ser His Asn
195 200 205
Glu Glu Ser Leu Gly Ser Thr Pro Pro Leu Pro Pro Gly Leu Pro Pro
210 215 220
Gly His Tyr Asp Ser Pro Lys Asn Ser His Ile Pro Gly His Tyr Asp
225 230 235 240
Leu Pro Pro Val Arg His Pro Pro Ser Pro Pro Ser Arg Arg Gln Asp
245 250 255
Arg

<210> 121
<211> 636
<212> PRT
<213> Rattus sp.

<400> 121
Met Gly Val Ile Cys Ser Leu Pro Cys Pro Glu Gly Phe His Gly Pro
1 5 10 15
Asn Cys Thr Gln Glu Cys Arg Cys His Asn Gly Gly Leu Cys Asp Arg
20 25 30
Phe Thr Gly Gln Cys His Cys Ala Pro Gly Tyr Ile Gly Asp Arg Cys
35 40 45
Arg Glu Glu Cys Pro Val Gly Arg Phe Gly Gln Asp Cys Ala Glu Thr
50 55 60

Cys	Asp	Cys	Ala	Pro	Gly	Ala	Arg	Cys	Phe	Pro	Ala	Asn	Gly	Ala	Cys
65					70				75					80	
Leu	Cys	Glu	His	Gly	Phe	Thr	Gly	Asp	Arg	Cys	Thr	Glu	Arg	Leu	Cys
				85					90					95	
Pro	Asp	Gly	Arg	Tyr	Gly	Leu	Ser	Cys	Gln	Asp	Pro	Cys	Thr	Cys	Asp
			100					105					110		
Pro	Glu	His	Ser	Leu	Ser	Cys	His	Pro	Met	His	Gly	Glu	Cys	Ser	Cys
		115					120					125			
Gln	Pro	Gly	Trp	Ala	Gly	Leu	His	Cys	Asn	Glu	Ser	Cys	Pro	Gln	Asp
	130					135					140				
Thr	His	Gly	Ala	Gly	Cys	Gln	Glu	His	Cys	Leu	Cys	Leu	His	Gly	Gly
145					150					155				160	
Val	Cys	Leu	Ala	Asp	Ser	Gly	Leu	Cys	Arg	Cys	Ala	Pro	Gly	Tyr	Thr
				165					170					175	
Gly	Pro	His	Cys	Ala	Asn	Leu	Cys	Pro	Pro	Asn	Thr	Tyr	Gly	Ile	Asn
			180					185					190		
Cys	Ser	Ser	His	Cys	Ser	Cys	Glu	Asn	Ala	Ile	Ala	Cys	Ser	Pro	Val
		195					200					205			
Asp	Gly	Thr	Cys	Ile	Cys	Lys	Glu	Gly	Trp	Gln	Arg	Gly	Asn	Cys	Ser
	210					215					220				
Val	Pro	Cys	Pro	Pro	Gly	Thr	Trp	Gly	Phe	Ser	Cys	Asn	Ala	Ser	Cys
225					230					235				240	
Gln	Cys	Ala	His	Glu	Gly	Val	Cys	Ser	Pro	Gln	Thr	Gly	Ala	Cys	Thr
				245					250					255	
Cys	Thr	Pro	Gly	Trp	Arg	Gly	Val	His	Cys	Gln	Leu	Pro	Cys	Pro	Lys
			260					265					270		
Gly	Gln	Phe	Gly	Glu	Gly	Cys	Ala	Ser	Val	Cys	Asp	Cys	Asp	His	Ser
		275				280						285			
Asp	Gly	Cys	Asp	Pro	Val	His	Gly	His	Cys	Arg	Cys	Gln	Ala	Gly	Trp
	290					295					300				
Met	Gly	Thr	Arg	Cys	His	Leu	Pro	Cys	Pro	Glu	Gly	Phe	Trp	Gly	Ala
305					310					315				320	
Asn	Cys	Ser	Asn	Ala	Cys	Thr	Cys	Lys	Asn	Gly	Gly	Thr	Cys	Val	Pro
			325						330					335	
Glu	Asn	Gly	Asn	Cys	Val	Cys	Ala	Pro	Gly	Phe	Arg	Gly	Pro	Ser	Cys
			340					345					350		
Gln	Arg	Pro	Cys	Pro	Pro	Gly	Arg	Tyr	Gly	Lys	Arg	Cys	Val	Pro	Cys
		355				360						365			
Lys	Cys	Asn	Asn	His	Ser	Ser	Cys	His	Pro	Ser	Asp	Gly	Thr	Cys	Ser
	370				375						380				
Cys	Leu	Ala	Gly	Trp	Thr	Gly	Pro	Asp	Cys	Ser	Glu	Ser	Cys	Pro	Pro
385					390					395				400	
Gly	His	Trp	Gly	Leu	Lys	Cys	Ser	Gln	Pro	Cys	Gln	Cys	His	His	Gly
			405						410					415	
Ala	Thr	Cys	His	Pro	Gln	Asp	Gly	Ser	Cys	Val	Cys	Ile	Pro	Gly	Trp
			420					425					430		
Thr	Gly	Pro	Asn	Cys	Ser	Glu	Gly	Cys	Pro	Ser	Arg	Met	Phe	Gly	Val
		435					440					445			
Asn	Cys	Ser	Gln	Leu	Cys	Gln	Cys	Asp	Pro	Gly	Glu	Met	Cys	His	Pro
	450					455					460				
Glu	Thr	Gly	Ala	Cys	Val	Cys	Pro	Pro	Gly	His	Ser	Gly	Ala	His	Cys
465					470					475				480	
Lys	Val	Gly	Ser	Gln	Glu	Ser	Phe	Thr	Ile	Met	Pro	Thr	Ser	Pro	Val
				485					490					495	
Ile	His	Asn	Ser	Leu	Gly	Ala	Val	Ile	Gly	Ile	Ala	Val	Leu	Gly	Thr
			500					505					510		
Leu	Val	Val	Ala	Leu	Val	Ala	Leu	Phe	Ile	Gly	Tyr	Arg	His	Trp	Gln
		515					520					525			

Lys Gly Lys Glu His Glu His Leu Ala Val Ala Tyr Ser Thr Gly Arg
 530 535 540
 Leu Asp Gly Ser Asp Tyr Val Met Pro Asp Val Ser Pro Ser Tyr Ser
 545 550 555 560
 His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser Gln Cys Ser Pro
 565 570 575
 Asn Pro Pro Pro Pro Asn Lys Ile Pro Gly Ser Gln Leu Phe Val Ser
 580 585 590
 Ser Gln Ala Ser Glu Arg Pro Asn Arg Asn His Gly Arg Asp Asn His
 595 600 605
 Ala Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu Ser His Asp Arg
 610 615 620
 Ala Phe Leu Arg His Gln Pro Pro Gly Pro Lys Val
 625 630 635

<210> 122
 <211> 500
 <212> PRT
 <213> Rattus sp.

<400> 122
 Met Gly Val Ile Cys Ser Leu Pro Cys Pro Glu Gly Phe His Gly Pro
 1 5 10 15
 Asn Cys Thr Gln Glu Cys Arg Cys His Asn Gly Gly Leu Cys Asp Arg
 20 25 30
 Phe Thr Gly Gln Cys His Cys Ala Pro Gly Tyr Ile Gly Asp Arg Cys
 35 40 45
 Arg Glu Glu Cys Pro Val Gly Arg Phe Gly Gln Asp Cys Ala Glu Thr
 50 55 60
 Cys Asp Cys Ala Pro Gly Ala Arg Cys Phe Pro Ala Asn Gly Ala Cys
 65 70 75 80
 Leu Cys Glu His Gly Phe Thr Gly Asp Arg Cys Thr Glu Arg Leu Cys
 85 90 95
 Pro Asp Gly Arg Tyr Gly Leu Ser Cys Gln Asp Pro Cys Thr Cys Asp
 100 105 110
 Pro Glu His Ser Leu Ser Cys His Pro Met His Gly Glu Cys Ser Cys
 115 120 125
 Gln Pro Gly Trp Ala Gly Leu His Cys Asn Glu Ser Cys Pro Gln Asp
 130 135 140
 Thr His Gly Ala Gly Cys Gln Glu His Cys Leu Cys Leu His Gly Gly
 145 150 155 160
 Val Cys Leu Ala Asp Ser Gly Leu Cys Arg Cys Ala Pro Gly Tyr Thr
 165 170 175
 Gly Pro His Cys Ala Asn Leu Cys Pro Pro Asn Thr Tyr Gly Ile Asn
 180 185 190
 Cys Ser Ser His Cys Ser Cys Glu Asn Ala Ile Ala Cys Ser Pro Val
 195 200 205
 Asp Gly Thr Cys Ile Cys Lys Glu Gly Trp Gln Arg Gly Asn Cys Ser
 210 215 220
 Val Pro Cys Pro Pro Gly Thr Trp Gly Phe Ser Cys Asn Ala Ser Cys
 225 230 235 240
 Gln Cys Ala His Glu Gly Val Cys Ser Pro Gln Thr Gly Ala Cys Thr
 245 250 255
 Cys Thr Pro Gly Trp Arg Gly Val His Cys Gln Leu Pro Cys Pro Lys
 260 265 270
 Gly Gln Phe Gly Glu Gly Cys Ala Ser Val Cys Asp Cys Asp His Ser
 275 280 285
 Asp Gly Cys Asp Pro Val His Gly His Cys Arg Cys Gln Ala Gly Trp

290 295 300
 Met Gly Thr Arg Cys His Leu Pro Cys Pro Glu Gly Phe Trp Gly Ala
 305 310 315 320
 Asn Cys Ser Asn Ala Cys Thr Cys Lys Asn Gly Gly Thr Cys Val Pro
 325 330 335
 Glu Asn Gly Asn Cys Val Cys Ala Pro Gly Phe Arg Gly Pro Ser Cys
 340 345 350
 Gln Arg Pro Cys Pro Pro Gly Arg Tyr Gly Lys Arg Cys Val Pro Cys
 355 360 365
 Lys Cys Asn Asn His Ser Ser Cys His Pro Ser Asp Gly Thr Cys Ser
 370 375 380
 Cys Leu Ala Gly Trp Thr Gly Pro Asp Cys Ser Glu Ser Cys Pro Pro
 385 390 395 400
 Gly His Trp Gly Leu Lys Cys Ser Gln Pro Cys Gln Cys His His Gly
 405 410 415
 Ala Thr Cys His Pro Gln Asp Gly Ser Cys Val Cys Ile Pro Gly Trp
 420 425 430
 Thr Gly Pro Asn Cys Ser Glu Gly Cys Pro Ser Arg Met Phe Gly Val
 435 440 445
 Asn Cys Ser Gln Leu Cys Gln Cys Asp Pro Gly Glu Met Cys His Pro
 450 455 460
 Glu Thr Gly Ala Cys Val Cys Pro Pro Gly His Ser Gly Ala His Cys
 465 470 475 480
 Lys Val Gly Ser Gln Glu Ser Phe Thr Ile Met Pro Thr Ser Pro Val
 485 490 495
 Ile His Asn Ser
 500

<210> 123
 <211> 24
 <212> PRT
 <213> Rattus sp.

<400> 123
 Leu Gly Ala Val Ile Gly Ile Ala Val Leu Gly Thr Leu Val Val Ala
 1 5 10 15
 Leu Val Ala Leu Phe Ile Gly Tyr
 20

<210> 124
 <211> 112
 <212> PRT
 <213> Rattus sp.

<400> 124
 Arg His Trp Gln Lys Gly Lys Glu His Glu His Leu Ala Val Ala Tyr
 1 5 10 15
 Ser Thr Gly Arg Leu Asp Gly Ser Asp Tyr Val Met Pro Asp Val Ser
 20 25 30
 Pro Ser Tyr Ser His Tyr Tyr Ser Asn Pro Ser Tyr His Thr Leu Ser
 35 40 45
 Gln Cys Ser Pro Asn Pro Pro Pro Asn Lys Ile Pro Gly Ser Gln
 50 55 60
 Leu Phe Val Ser Ser Gln Ala Ser Glu Arg Pro Asn Arg Asn His Gly
 65 70 75 80
 Arg Asp Asn His Ala Thr Leu Pro Ala Asp Trp Lys His Arg Arg Glu
 85 90 95
 Ser His Asp Arg Ala Phe Leu Arg His Gln Pro Pro Gly Pro Lys Val

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100
105
110

<210> 125
<211> 28
<212> PRT
<213> Homo sapiens

<400> 125
Met Ala Pro Ala Arg Ala Gly Phe Cys Pro Leu Leu Leu Leu Leu Leu
1      5      10      15
Leu Gly Leu Trp Val Ala Glu Ile Pro Val Ser Ala
20      25

<210> 126
<211> 128
<212> PRT
<213> Homo sapiens

<400> 126
Lys Pro Lys Gly Met Thr Ser Ser Gln Trp Phe Lys Ile Gln His Met
1      5      10      15
Gln Pro Ser Pro Gln Ala Cys Asn Ser Ala Met Lys Asn Ile Asn Lys
20      25      30
His Thr Lys Arg Cys Lys Asp Leu Asn Thr Phe Leu His Glu Pro Phe
35      40      45
Ser Ser Val Ala Ala Thr Cys Gln Thr Pro Lys Ile Ala Cys Lys Asn
50      55      60
Gly Asp Lys Asn Cys His Gln Ser His Gly Pro Val Ser Leu Thr Met
65      70      75      80
Cys Lys Leu Thr Ser Gly Lys Tyr Pro Asn Cys Arg Tyr Lys Glu Lys
85      90      95
Arg Gln Asn Lys Ser Tyr Val Val Ala Cys Lys Pro Pro Gln Lys Lys
100      105      110
Asp Ser Gln Gln Phe His Leu Val Pro Val His Leu Asp Arg Val Leu
115      120      125

<210> 127
<211> 19
<212> PRT
<213> Homo sapiens

<400> 127
Met Pro Leu Leu Thr Leu Tyr Leu Leu Leu Phe Trp Leu Ser Gly Tyr
1      5      10      15
Ser Ile Ala

<210> 128
<211> 286
<212> PRT
<213> Homo sapiens

<400> 128
Thr Gln Ile Thr Gly Pro Thr Thr Val Asn Gly Leu Glu Arg Gly Ser
1      5      10      15
Leu Thr Val Gln Cys Val Tyr Arg Ser Gly Trp Glu Thr Tyr Leu Lys
20      25      30
Trp Trp Cys Arg Gly Ala Ile Trp Arg Asp Cys Lys Ile Leu Val Lys

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<210> 130
 <211> 24
 <212> PRT
 <213> Homo sapiens

<400> 130
 Leu Ser Val Leu Leu Pro Leu Ile Phe Thr Ile Leu Leu Leu Leu Leu
 1 5 10 15
 Val Ala Ala Ser Leu Leu Ala Trp
 20

<210> 131
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 131
 Arg Met Met Lys Tyr Gln Gln Lys Ala Ala Gly Met Ser Pro Glu Gln
 1 5 10 15
 Val Leu Gln Pro Leu Glu Gly Asp Leu Cys Tyr Ala Asp Leu Thr Leu
 20 25 30
 Gln Leu Ala Gly Thr Ser Pro Arg Lys Ala Thr Thr Lys Leu Ser Ser
 35 40 45
 Ala Gln Val Asp Gln Val Glu Val Glu Tyr Val Thr Met Ala Ser Leu
 50 55 60
 Pro Lys Glu Asp Ile Ser Tyr Ala Ser Leu Thr Leu Gly Ala Glu Asp
 65 70 75 80
 Gln Glu Pro Thr Tyr Cys Asn Met Gly His Leu Ser Ser His Leu Pro
 85 90 95
 Gly Arg Gly Pro Glu Glu Pro Thr Glu Tyr Ser Thr Ile Ser Arg Pro
 100 105 110

<210> 132
 <211> 21
 <212> PRT
 <213> Homo sapiens

<400> 132
 Met Asp His Cys Gly Ala Leu Phe Leu Cys Leu Cys Leu Leu Thr Leu
 1 5 10 15
 Gln Asn Ala Thr Thr
 20

<210> 133
 <211> 507
 <212> PRT
 <213> Homo sapiens

<400> 133
 Glu Thr Trp Glu Glu Leu Leu Ser Tyr Met Glu Asn Met Gln Val Ser
 1 5 10 15
 Arg Gly Arg Ser Ser Val Phe Ser Ser Arg Gln Leu His Gln Leu Glu
 20 25 30
 Gln Met Leu Leu Asn Thr Ser Phe Pro Gly Tyr Asn Leu Thr Leu Gln
 35 40 45
 Thr Pro Thr Ile Gln Ser Leu Ala Phe Lys Leu Ser Cys Asp Phe Ser
 50 55 60
 Gly Leu Ser Leu Thr Ser Ala Thr Leu Lys Arg Val Pro Gln Ala Gly

65					70					75				80
Gly	Gln	His	Ala	Arg	Gly	Gln	His	Ala	Met	Gln	Phe	Pro	Ala	Glu
				85					90					95
Thr	Arg	Asp	Ala	Cys	Lys	Thr	Arg	Pro	Arg	Glu	Leu	Arg	Leu	Ile
			100					105						110
Ile	Tyr	Phe	Ser	Asn	Thr	His	Phe	Phe	Lys	Asp	Glu	Asn	Asn	Ser
		115					120					125		
Leu	Leu	Asn	Asn	Tyr	Val	Leu	Gly	Ala	Gln	Leu	Ser	His	Gly	His
	130						135				140			Val
Asn	Asn	Leu	Arg	Asp	Pro	Val	Asn	Ile	Ser	Phe	Trp	His	Asn	Gln
145				150						155				160
Leu	Glu	Gly	Tyr	Thr	Leu	Thr	Cys	Val	Phe	Trp	Lys	Glu	Gly	Ala
				165					170					175
Lys	Gln	Pro	Trp	Gly	Gly	Trp	Ser	Pro	Glu	Gly	Cys	Arg	Thr	Glu
		180						185						190
Pro	Ser	His	Ser	Gln	Val	Leu	Cys	Arg	Cys	Asn	His	Leu	Thr	Tyr
	195						200					205		Phe
Ala	Val	Leu	Met	Gln	Leu	Ser	Pro	Ala	Leu	Val	Pro	Ala	Glu	Leu
	210						215				220			Leu
Ala	Pro	Leu	Thr	Tyr	Ile	Ser	Leu	Val	Gly	Cys	Ser	Ile	Ser	Ile
225					230					235				240
Ala	Ser	Leu	Ile	Thr	Val	Leu	Leu	His	Phe	His	Phe	Arg	Lys	Gln
				245					250					255
Asp	Ser	Leu	Thr	Arg	Ile	His	Met	Asn	Leu	His	Ala	Ser	Val	Leu
	260							265					270	Leu
Leu	Asn	Ile	Ala	Phe	Leu	Leu	Ser	Pro	Ala	Phe	Ala	Met	Ser	Pro
	275						280					285		Val
Pro	Gly	Ser	Ala	Cys	Thr	Ala	Leu	Ala	Ala	Ala	Leu	His	Tyr	Ala
	290					295					300			Leu
Leu	Ser	Cys	Leu	Thr	Trp	Met	Ala	Ile	Glu	Gly	Phe	Asn	Leu	Tyr
305					310					315				320
Leu	Leu	Gly	Arg	Val	Tyr	Asn	Ile	Tyr	Ile	Arg	Arg	Tyr	Val	Phe
				325					330					335
Leu	Gly	Val	Leu	Gly	Trp	Gly	Ala	Pro	Ala	Leu	Leu	Val	Leu	Leu
		340					345					350		Ser
Leu	Ser	Val	Lys	Ser	Ser	Val	Tyr	Gly	Pro	Cys	Thr	Ile	Pro	Val
	355					360						365		Phe
Asp	Ser	Trp	Glu	Asn	Gly	Thr	Gly	Phe	Gln	Asn	Met	Ser	Ile	Cys
	370					375					380			Trp
Val	Arg	Ser	Pro	Val	Val	His	Ser	Val	Leu	Val	Met	Gly	Tyr	Gly
385					390					395				400
Leu	Thr	Ser	Leu	Phe	Asn	Leu	Val	Val	Leu	Ala	Trp	Ala	Leu	Trp
				405					410					415
Leu	Arg	Arg	Leu	Arg	Glu	Arg	Ala	Asp	Ala	Pro	Ser	Val	Arg	Ala
				420				425					430	Cys
His	Asp	Thr	Val	Thr	Val	Leu	Gly	Leu	Thr	Val	Leu	Leu	Gly	Thr
	435						440					445		Thr
Trp	Ala	Leu	Ala	Phe	Phe	Ser	Phe	Gly	Val	Phe	Leu	Leu	Pro	Gln
	450					455				460				Leu
Phe	Leu	Phe	Thr	Ile	Leu	Asn	Ser	Leu	Tyr	Gly	Phe	Phe	Leu	Phe
465					470					475				480
Trp	Phe	Cys	Ser	Gln	Arg	Cys	Arg	Ser	Glu	Ala	Glu	Ala	Lys	Ala
				485					490					495
Ile	Glu	Ala	Phe	Ser	Ser	Ser	Gln	Thr	Gln					
			500					505						

<210> 134
<211> 223

<212> PRT
 <213> Homo sapiens

<400> 134
 Glu Thr Trp Glu Glu Leu Leu Ser Tyr Met Glu Asn Met Gln Val Ser
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 Arg Gly Arg Ser Ser Val Phe Ser Ser Arg Gln Leu His Gln Leu Glu
 20 25 30
 Gln Met Leu Leu Asn Thr Ser Phe Pro Gly Tyr Asn Leu Thr Leu Gln
 35 40 45
 Thr Pro Thr Ile Gln Ser Leu Ala Phe Lys Leu Ser Cys Asp Phe Ser
 50 55 60
 Gly Leu Ser Leu Thr Ser Ala Thr Leu Lys Arg Val Pro Gln Ala Gly
 65 70 75 80
 Gly Gln His Ala Arg Gly Gln His Ala Met Gln Phe Pro Ala Glu Leu
 85 90 95
 Thr Arg Asp Ala Cys Lys Thr Arg Pro Arg Glu Leu Arg Leu Ile Cys
 100 105 110
 Ile Tyr Phe Ser Asn Thr His Phe Lys Asp Glu Asn Asn Ser Ser
 115 120 125
 Leu Leu Asn Asn Tyr Val Leu Gly Ala Gln Leu Ser His Gly His Val
 130 135 140
 Asn Asn Leu Arg Asp Pro Val Asn Ile Ser Phe Trp His Asn Gln Ser
 145 150 155 160
 Leu Glu Gly Tyr Thr Leu Thr Cys Val Phe Trp Lys Glu Gly Ala Arg
 165 170 175
 Lys Gln Pro Trp Gly Gly Trp Ser Pro Glu Gly Cys Arg Thr Glu Gln
 180 185 190
 Pro Ser His Ser Gln Val Leu Cys Arg Cys Asn His Leu Thr Tyr Phe
 195 200 205
 Ala Val Leu Met Gln Leu Ser Pro Ala Leu Val Pro Ala Glu Leu
 210 215 220

<210> 135
 <211> 25
 <212> PRT
 <213> Homo sapiens

<400> 135
 Leu Ala Pro Leu Thr Tyr Ile Ser Leu Val Gly Cys Ser Ile Ser Ile
 1 5 10 15
 Val Ala Ser Leu Ile Thr Val Leu Leu
 20 25

<210> 136
 <211> 20
 <212> PRT
 <213> Homo sapiens

<400> 136
 Leu His Ala Ser Val Leu Leu Leu Asn Ile Ala Phe Leu Leu Ser Pro
 1 5 10 15
 Ala Phe Ala Met
 20

<210> 137
 <211> 21
 <212> PRT

<213> Homo sapiens

<400> 137

Tyr Ala Leu Leu Ser Cys Leu Thr Trp Met Ala Ile Glu Gly Phe Asn
1 5 10 15
Leu Tyr Leu Leu Leu
20

<210> 138

<211> 19

<212> PRT

<213> Homo sapiens

<400> 138

Leu Gly Val Leu Gly Trp Gly Ala Pro Ala Leu Leu Val Leu Leu Ser
1 5 10 15
Leu Ser Val

<210> 139

<211> 25

<212> PRT

<213> Homo sapiens

<400> 139

Val Leu Val Met Gly Tyr Gly Gly Leu Thr Ser Leu Phe Asn Leu Val
1 5 10 15
Val Leu Ala Trp Ala Leu Trp Thr Leu
20 25

<210> 140

<211> 21

<212> PRT

<213> Homo sapiens

<400> 140

Val Thr Val Leu Gly Leu Thr Val Leu Leu Gly Thr Thr Trp Ala Leu
1 5 10 15
Ala Phe Phe Ser Phe
20

<210> 141

<211> 20

<212> PRT

<213> Homo sapiens

<400> 141

Leu Phe Leu Phe Thr Ile Leu Asn Ser Leu Tyr Gly Phe Phe Leu Phe
1 5 10 15
Leu Trp Phe Cys
20

<210> 142

<211> 24

<212> PRT

<213> Homo sapiens

<400> 142

Ser Gln Arg Cys Arg Ser Glu Ala Glu Ala Lys Ala Gln Ile Glu Ala
 1 5 10 15
 Phe Ser Ser Ser Gln Thr Thr Gln
 20

<210> 143
 <211> 16
 <212> PRT
 <213> Homo sapiens

<400> 143
 Ser Pro Val Pro Gly Ser Ala Cys Thr Ala Leu Ala Ala Ala Leu His
 1 5 10 15

<210> 144
 <211> 37
 <212> PRT
 <213> Homo sapiens

<400> 144
 Lys Ser Ser Val Tyr Gly Pro Cys Thr Ile Pro Val Phe Asp Ser Trp
 1 5 10 15
 Glu Asn Gly Thr Gly Phe Gln Asn Met Ser Ile Cys Trp Val Arg Ser
 20 25 30
 Pro Val Val His Ser
 35

<210> 145
 <211> 7
 <212> PRT
 <213> Homo sapiens

<400> 145
 Gly Val Phe Leu Leu Pro Gln
 1 5

<210> 146
 <211> 17
 <212> PRT
 <213> Homo sapiens

<400> 146
 His Phe His Phe Arg Lys Gln Ser Asp Ser Leu Thr Arg Ile His Met
 1 5 10 15
 Asn

<210> 147
 <211> 14
 <212> PRT
 <213> Homo sapiens

<400> 147
 Gly Arg Val Tyr Asn Ile Tyr Ile Arg Arg Tyr Val Phe Lys
 1 5 10

<210> 148
 <211> 18

<212> PRT

<213> Homo sapiens

<400> 148

Arg	Arg	Leu	Arg	Glu	Arg	Ala	Asp	Ala	Pro	Ser	Val	Arg	Ala	Cys	His
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Asp	Thr														

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C07K 14/47; C07H 21/04; C12N 15/68; C12P 21/08

US CL : 530/550; 535/55.5; 435/520.1, 552.9, 561, 59.1

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 530/550; 535/55.5; 435/520.1, 552.9, 561, 59.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

Commercial Sequence Databases: GenEmbl, EST, Issued_Patents_NA, N_Geneseq_98, PIR_94, SwissProt_98, A_Geneseq_98, Issued_Patents_AA, SPTREMBL_19

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Database EST, AN AQ588144, ZHOU et al. 'CITBI-E1-2644L24.TF CITBI-E1 Homo sapiens genomic clone 2644L24, genomic survey sequence'. 07 June 1999, see attached alignment showing 100% identical match to nucleotides 88-481 of SEQ ID NO: 1 (394 nucleotides total).	1, 3, 5
Y		2, 4, 6-10 and 12
A	Database SPTREMBL_12, AN Q28396, RICHARDSON et al. 'Type II Collagen from Equus caballus (Horse)'. 01 November 1996. Polypeptide 25.7% identical to the amino acid sequence of SEQ ID NO:2, see attached alignment, Nov. 1, 1996.	1-10 and 12

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	* Later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
* A Document defining the general state of the art which is not considered to be of particular relevance	* X Document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
* F Earlier document published on or after the international filing date	* Y Document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other cited documents, such combination being obvious to a person skilled in the art
* I Document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	* Z Document member of the same patent family
* O Document referring to an oral disclosure, use, exhibition or other means	
* P Document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

21 SEPTEMBER 2000

Date of mailing of the international search report

02 OCT 2000

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
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Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-10 and 12

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (continuation of first sheet(1)) (July 1998)*

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING

This ISA found multiple inventions as follows:

This application contains the following inventions or groups of inventions which are not so linked as to form a single inventive concept under PCT Rule 13.1. In order for all inventions to be searched, the appropriate additional search fees must be paid.

Group I, claim(s) 1-10 and 12, in so far as they are drawn to Intercept 340, polynucleotides of SEQ ID NOS: 1 and 3, vector, host cell, method of producing a protein recombinantly and protein of SEQ ID NO: 2.

Groups II-VII, claim(s) 1-10 and 12, in so far as they are drawn to the next six polynucleotides of distinct cDNA clones and encoded proteins, identified as Mango 003, Mango 347, Tango 272, Tango 295, Tango 354 and Tango 378, as listed in Tables 1 and 2.

Groups VIII-XIV, claim(s) 11 and 15, in so far as they are drawn to antibodies to one of the seven proteins listed above.

Groups XV-XXI, claims 13, 14, 19, 20 and 22, in so far as they are drawn to a method for detecting the presence of in a sample or identifying a compound which binds to or modulates the activity of a polypeptide of one of the seven proteins listed above.

Groups XXII-XXVII, claims 16 and 17, in so far as they are drawn to a method for detecting the nucleic acids of one of the seven cDNA clones listed above.

Groups XXIX-XXXV, claim 18, in so far as it is drawn to a kit comprising a compound of unspecified constitution which selectively binds to a nucleic acid molecule of the seven cDNA clones listed above.

Groups XXXVI-XLII, claim 21, in so far as it is drawn to a method for modulating the activity of one of the seven proteins listed above.

The inventions listed as Groups I-XLII do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Group I corresponds to the first invention wherein the first product is the polynucleotide and the first method of using is the method of making the protein. Note that there is no method of making the polynucleotide. The invention also includes the protein made. Each of groups II-VII does not share the same or corresponding special technical feature because each group is drawn to a different polynucleotide and encoded protein, and each of groups VIII-XLII does not share the same or corresponding special technical feature because each group is drawn to different compounds or methods of using the seven polynucleotides and encoded proteins. This Authority therefore considers that the several inventions do not share a special technical feature within the meaning of PCT Rule 13.2 and thus do not relate to a single general inventive concept within the meaning of PCT Rule 13.1.